



Europäisches Patentamt
European Patent Office
Office européen des brevets

(11) Publication number:

0 097 373

A2

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 83105112.2

(61) Int. Cl.³: C 07 H 21/00

(22) Date of filing: 22.06.83

C 07 H 19/04, C 07 G 7/00
A 61 K 31/70, C 12 P 21/00
G 01 N 33/48, C 07 C 155/03

(30) Priority: 23.06.82 US 391440

(72) Inventor: Rabbani, Elazar
69 Fifth Avenue
New York, N.Y. 10003(US)

(43) Date of publication of application:
04.01.84 Bulletin 84/1

(72) Inventor: Kline, Stanley
235 Lincoln Place
Brooklyn, N.Y. 11217(US)

(84) Designated Contracting States:
AT BE CH DE FR GB IT LI LU NL SE

(72) Inventor: Stavrianopoulos, Jannis G.
515 West 59 Street
New York, N.Y. 10019(US)

(71) Applicant: ENZO BIOCHEM, INC.
325 Hudson Street
New York, N.Y. 10013(US)

(72) Inventor: Kirtikar, Dollie
42-72 80 Street
Elmhurst, N.Y. 11373(US)

(72) Inventor: Engelhardt, Dean
173 Riverside Drive
New York, N.Y. 10024(US)

(72) Representative: Vossius Vossius Tauchner Heunemann
Rauh
Siabertstrasse 4 P.O. Box 86 07 67
D-8000 München 86(DE)

(54) Modified labeled nucleotides and polynucleotides and methods of preparing, utilizing and detecting same.

(57) Nucleotides and polynucleotides, including DNA, are chemically modified or labeled so as to be capable of ready detection when attached to and/or incorporated in nucleic acid material.

0097373

1 Our Ref.: S 458EP
Case: 19439

ENZO BIOCHEM, INC.
New York, N.Y.

MODIFIED LABELED NUCLEOTIDES AND POLYNUCLEOTIDES AND
METHODS OF PREPARING, UTILIZING AND DETECTING SAME
BACKGROUND OF THE INVENTION

5

It is known to produce nucleotides or polynucleotides which are radioactively labeled, such as with isotopes or hydrogen (³H), phosphorus (³²P), carbon (¹⁴C) or iodine (¹²⁵I). Such radioactively labeled compounds are 10 useful to detect, monitor, localize and isolate nucleic acids and other molecules of scientific or clinical interest. Unfortunately, however, the use of radioactively labeled materials presents hazards due to radiation. Also due to the relatively short half life 15 of the radioactive materials employed to label such compounds or materials, the resulting labeled compounds or materials have a corresponding relatively short shelf life.

20 It has been proposed to chemically label compounds of interest, such as nucleotides and polynucleotides, so as to overcome or avoid the hazards and difficulties associated with such compounds or materials when radioactively labeled. In the article by P.R. Langer,

25 A. A. Waldrop and D. C. Ward entitled "Enzymatic Synthesis of Biotin-Labeled Polynucleotides: Novel Nucleic Acid Affinity Probes", in Proc. Natl. Acad. Sci., USA, Vol. 78, No. 11, pp. 6633-6637, November,

30 1981, there are described analogs of dUTP and UTP that contain a biotin molecule bound to the C-5 position of the pyrimidine ring through an alkylamine linker arm. The biotin-labeled nucleotides are efficient substrates for a variety of DNA and RNA polymerases in vitro.

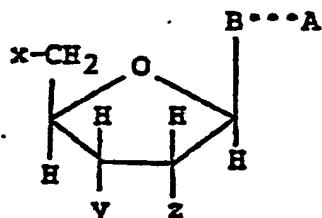
35 Biotin-labeled polynucleotides containing low levels of biotin substitution (50 molecules or fewer per kilobase) have denaturation, reassociation and hybridization characteristics similar to those of unsubstituted controls. Biotin-labeled polynucleotides, both single and double

0097373

1 stranded, are selectively and quantitatively retained on
avidin-Sephar se, even after extensive washing with 8M
urea, 6M guanidine hydrochloride or 99% formamide. In
addition, biotin-labeled nucleotides can be selectively
5 immunoprecipitated in the presence of antibiotin
antibody and Staphylococcus aurea, Protein A. These
unique features of biotin-labeled polynucleotides
suggest that they are useful affinity probes for the
detection and isolation of specific DNA and RNA
10 sequences.

The disclosures of this article are herein incorporated
and made part of this disclosure. The subject matter of
15 said article is comprised in EP-A2-0063879 in which
additionally it is disclosed that
compounds having the structure:

20



25

30

35.

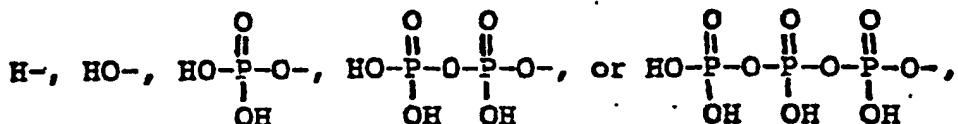
0097373

1 wherein B represents a purine, deazapurine, or pyrimidine moiety covalently bonded to the C¹-position of the sugar moiety, provided that when B is purine or 7-deazapurine, it is attached at the N⁹-position of the purine or deazapurine,
5 and when B is pyrimidine, it is attached at the N¹-position;

wherein A represents a moiety consisting of at least three carbon atoms which is capable of forming a detectable complex with a polypeptide when the compound is incorporated
10 into a double-stranded ribonucleic acid, deoxyribonucleic acid duplex, or DNA-RNA hybrid;

wherein the dotted line represents a chemical linkage joining B and A, provided that if B is purine the linkage is
15 attached to the 8-position of the purine, if B is 7-deazapurine, the linkage is attached to the 7-position of the deazapurine, and if B is pyrimidine, the linkage is attached to the 5-position of the pyrimidine; and

20 wherein each of x, y, and z represents



25 either directly, or when incorporated into oligo- and polynucleotides, provide probes which are widely usely.

Applications disclosed in EP-A2-0 063 879 include detection and localization of polynucleotide sequences in chromosomes, fixed cells, tissue sections and cell extracts.
80 Specific applications include chromosomal karyotyping, clinical diagnosis of nucleic acid-containing etiological agents, e.g. bacteria, viruses, or fungi, and diagnosis of genetic disorders.

35

The disclosures of EP-A2-0 063 879 are herein incorporated and made part of this disclosure.

By way of additional background with respect to the utilization of the biotin-polynucleotides of the above-identified Langer et al article in Proc. Natl. Acad. Sci. USA, the publication by P. R. Langer-Safer, M. 05 Levine and D. C. Ward in Genetics entitled "An Immunological Method for Mapping Genes on Drosophila Polytene Chromosomes", describes a method employing biotinylated nucleotides as a probe for the localization of DNA sequences hybridized *in situ* to Drosophila 10 polytene chromosomes. In this application these probes are detected using affinity purified rabbit antibiotin antibody as the primary antibody and fluorescinated goat antirabbit antibody as the secondary antibody. The disclosures of this Langer-Safer et al publication in 15 Genetics are also incorporated and made part of this disclosure.

Other techniques employing biotin-labeled reagents with avidin or enzyme-labeled avidin reagents are known for 20 the detection and determination of ligands in a liquid medium, see U.S. Patent 4,228,237. Also, it is known to effect gene enrichment based on avidin-biotin interaction, particularly as applied to Drosophila ribosomal RNA genes, see the J. Manning, M. Pellegrini and N. 25 Davidson publication in Biochemistry, Vol. 16, No. 7, pages 1364-1369 (1977). Other publications of background interest with respect to the practices of this invention are the D. J. Eckermann and R. H. Symons article entitled "Sequence at the Site of Attachment of 30 an Affinity-Label Derivative of Puromycin on 23-S Ribosomal RNA of Escherichia coli Ribosomes", J. Biochem., 82, 225-234 (1978); the article by S. B. Zimmerman, S. R. Kornberg and A. Kornberg entitled "Glucosylation of Deoxyribonucleic Acid-II -Glucosyl Transferases from 35 T2- and T6-Infected Escherichia coli in The Journal of Biological Chemistry", Vol. 237, No. 2, February 1962,

and the article by J. Josse and A. Kornberg "III. α - and β -Glucosyl Transferases from T4-Infected Escherichia coli", also appearing in The Journal of Biological Chemistry, Vol. 237, No. 6, June 1962.

- 05 Of further interest in connection with the practices of this invention are the publications appearing in the J. Biol. Chem., Vol. 236, No. 5, May 1961, pages 1487-1493; the same publication, Vol. 237, No. 4, pages 1251-1259 (1962); the same publication Vol. 239, No. 9, 10 pages 2957-2963 (1964). Of special interest is the article appearing The Journal of Histochemistry and Cytochemistry, Vol. 27, No. 8, pages 1131-1139 (1979) and in the publication Nucleic Acids Research, Vol. 5, No. 9, 1977, pages 2961-2973. Also of interest is the 15 article appearing in the publication Biochimica et Biophysica Acta by A. De Waard entitled "Specificity Difference Between the Hydroxymethylcytosine β -Glucosyl-Transferases Induced by Bacteriophages T2, T4 and T6", pages 286-304, and also the article by T. W. North and 20 C. K. Mathews entitled "T4 Phage-Coded Deoxycytidylate Hydroxymethylase: Purification and Studies in Inter-molecular Interactions", published by Academic Press, 1977, pages 898-904 and the article by E. A. Bayer and M. Wilchek entitled "The Use of Avidin-Biotin Complex as 25 a Tool in Molecular Biology in Methods of Biochemical Analysis, Vol. 26, pages 1-45 (1980).

Other techniques useful in the practices of this invention include nick translation of DNA employing DNA polymerase. A technique for effecting nick translation is disclosed in the article by P. W. Rigby, M. Dieckmann, 05 C. Rhodes and P. Berg entitled "Labeling Deoxyribonucleic Acid to High Specific Activity in vitro by Nick Translation with DNA Polymerase" in J. Mol. Biol. (1977), 113, 237-251. With respect to the recovery of streptavidin, such as from a culture broth of Streptomyces avidinii, 10 the article by K. Hofmann, S. W. Wood, C. C. Brinton, J. A. Montibeller and F. M. Finn entitled "Iminobiotin Affinity Columns and their Application to Retrieval of Streptavidin" in Proc. Natl. Acad. Sci. USA, Vol. 77, No. 8, pp. 4666-4668 (1980), discloses a suitable 15 approach for the recovery of streptavidin from a streptavidin-containing material, such as from a culture broth. Streptavidin is useful as a reagent in one of the practices of this invention.

20 The aforementioned publications are herein incorporated and made part of this disclosure.

SUMMARY OF THE INVENTION

25 In accordance with the practices of this invention nucleotides are modified, such as at the 5 position of pyrimidine or the 7 position of purine, preparatory for the preparation therefrom of nucleotide probes suitable for attachment to or incorporation into DNA or other 30 nucleic acid material. In the practices of this invention nucleotides, i.e. nucleic acids, preferably are modified in a non-disruptive manner such that the resulting modified nucleotides are capable of incorporation into nucleic acids and once incorporated in

nucleic acids the modified nucleotides do not significantly interfere with the formation or stabilization of the double helix formed of the resulting nucleic acids containing the modified nucleotides. The non-disruptive
05 modification of nucleotides and nucleic acids incorporating such modified nucleotides is in contrast with those modifications of nucleotides which are characterized as a disruptive modification in the sense that the resulting disruptively modified nucleotides and nucleic
10 acids containing the same block proper double helix formation. In the practices of this invention, the nucleotides are desirably modified at the 5 position of the pyrimidine or the 7 position of the purine. The nucleotides so modified are non-disruptively modified
15 and nucleic acids containing such nucleotides are capable of forming a double helix arrangement..

Broadly, in another aspect of the practices of this invention various methods are useful for the tagging or
20 labeling of DNA in a non-disruptive manner. For example, biotin is added on the end of a DNA or RNA molecule. The addition of biotin is accomplished by addition of a ribonucleotide. The 3',4' vicinal hydroxyl groups are oxidized by periodate oxidation and
25 then reduced by a borohydride in the presence of biotin hydrazide. Alternatively, carbodiimide can also be used to couple biotin to the aldehyde group.

Another technique for tagging nucleic acid material such as DNA or RNA involves the addition of a large marker to
30 the end of a DNA or RNA molecule. One example of this technique is the addition of a molecule, e.g. lysyl-glycine, where the amino groups are tagged with biotin. Another example would be to follow the procedure set
35 forth hereinabove but employing carbodiimide as the

cross-linking agent. Still another example of this technique would be to produce a biotinylated dA:dV double helical polymer and to ligate this polymer to the probe prepared in accordance with this invention.

05

Another technique for tagging DNA in a non-disruptive manner involves the isolation of dPyrTP having a putricine or spermidine on the 5 position from PS16 or phage-infected cells. If desired, dPyrTP is made from 10 phage DNA and phosphorylated to dPyrTP followed by modification of the polyamine side chain by means of standard nucleophilic reagent NHS-biotin.

15 Another technique for tagging DNA in a non-disruptive manner involves the addition of glucose to 5-hydroxy-methylcytosine (5 HMC) in DNA using T4 phage glycoslyating enzymes followed by screening by means of a lectin-based assay.

20 Still another method for tagging DNA in a non-disruptive manner involves 5-HMC-triphosphate made from the hydrolysis of T4-DNA followed by phosphorylation of the 5HMCMMP to 5 HMCTP. 5HMCTP is then incorporated into DNA using polymerase I. Thus, any DNA can be modified to 25 have non-disruptively incorporated therein 5 HMC.

30 A method for tagging DNA in a mildly disruptive manner involves reacting nucleic acids in the double helical form with alkylating reagents as for example benz(o)pyrene diol epoxide or aflatoxin. Under appropriate conditions the N² group of guanine, the N⁴ group of adenosine or the N⁴ group of cytosine are alkylated. These modified nucleotides can be directly detected with 35 antibodies or can be used as linking arms for the addition of a reporter molecule such as biotin.

1 Preferred embodiments of the present invention are explained in detail in the following enumeration.

5 1. A nucleotide having the general formula P-S-B-Sig wherein P is the phosphoric acid moiety, S the sugar or monosaccharide moiety, B being the base moiety, the phosphoric acid moiety being attached at the 3' and/or the 5' position of the sugar moiety when said nucleotide
10 is a deoxyribonucleotide and at the 2', 3' and/or 5' position when said nucleotide is a ribonucleotide, said base being a purine or a pyrimidine, said base being attached from the N1 position or the N9 position to the 1' position of the sugar moiety when said base is a
15 pyrimidine or a purine, respectively, and wherein said Sig is a chemical moiety covalently attached to the base B of said nucleotide, said Sig when attached to said base B being capable of signalling itself or makes itself self-detecting or its presence known.

20 2. A nucleotide in accordance with item 1 wherein said nucleotide is a deoxyribonucleotide.

25 3. A nucleotide in accordance with item 1 wherein said nucleotide is a ribonucleotide.

30 4. A nucleotide in accordance with item 1 wherein said chemical moiety Sig is chemically attached to B at the N7 position when B is a 7-deazapurine, at the C5 position when B is a pyrimidine and at the C8 position when B is a purine.

- 1 5. A nucleotide in accordance with item 1 wherein Sig is attached to B at a position such that an oligonucleotide or polynucleotide containing said nucleotide is capable of forming a double-stranded ribonucleic acid, a
- 5 double-stranded deoxyribonucleic acid or a DNA-RNA hybrid, or when said nucleotide is incorporated into said oligonucleotide or polynucleotide.
- 10 6. A nucleotide in accordance with item 1 wherein Sig is attached to B at a position such that said nucleotide is capable of being incorporated into or to form a double-stranded ribonucleic acid, a double-stranded deoxyribonucleic acid or a double-stranded deoxyribonucleic acid-ribonucleic acid hybrid.
- 15 7. A nucleotide in accordance with item 1 wherein Sig is attached to B at a position such that when said nucleotide is incorporated into or attached to or associated with a double-stranded deoxyribonucleic acid or double-stranded ribonucleic acid or DNA-RNA hybrid, said chemical moiety Sig is capable of signalling itself or making itself self-detecting or its presence known.
- 20 8. An oligonucleotide or polydeoxyribonucleotide comprising one or more nucleotides in accordance with item 1.
- 25 9. An oligonucleotide or polyribonucleotide comprising one or more nucleotides in accordance with item 1.

- 1 10. A nucleotide in accordance with item 1 wherein said Sig is attached to said base B at the C5 position when said base B is a pyrimidine or at the C7 position when said base B is a deazapurin.
- 5 11. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is an aliphatic chemical moiety containing at least 4 carbon atoms.
- 10 12. A nucleotide in accordance with item 1 wherein said base B is a pyrimidine and wherein said Sig chemical moiety is attached to the pyrimidine at the N3 position.
- 15 13. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is an aliphatic chemical moiety containing at least 3 carbon atoms and at least one double bond.
- 20 14. A nucleotide in accordance with item 1 wherein said base B is a pyrimidine and wherein said Sig chemical moiety is attached to the pyrimidine at the C5 position.
- 25 15. A nucleotide in accordance with item 1 wherein said base B is a pyrimidine and wherein said Sig chemical moiety is attached to the pyrimidine at the C6 position.
- 30 16. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety is attached to the purine at the N1 position.

17. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety is attached to the purine at the C2 position.

05 18. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety is attached to the purine at the N3 position.

10 19. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety contains an aromatic or a cycloaliphatic group containing at least six or at least five carbon atoms, respectively.

15 20. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety is attached to the purine at the N7 position.

20 21. A nucleotide in accordance with item 1 wherein said base B is a purine and wherein said Sig chemical moiety is attached to the purine at the C8 position.

25 22. A nucleotide in accordance with item 1 wherein said S sugar is a pentose.

23. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is a polysaccharide or an oligosaccharide.

30 24. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is a sugar selected from the group consisting of triose or tetrose, or pentose, a hexose, or heptose, and an octose.

25. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is attached by or includes a glycosidic linkage moiety.

05 26. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is a hexose moiety.

10 27. A nucleotide in accordance with item 1 wherein said base B is a 7-deazapurine and wherein said Sig chemical moiety is attached to the 7-deazapurine at the C7 position.

15 28. A nucleotide in accordance with item 1 wherein said Sig chemical moiety comprises a component selected from the group consisting of an electron dense component, a magnetic component, an enzyme, a hormone component, a radioactive component, a metal-containing component, a fluorescing component and an antigen or antibody component.

20 29. A nucleotide in accordance with item 1 wherein said Sig chemical moiety is a sugar residue and wherein said sugar is complexed with or attached to a sugar or polysaccharide binding protein.

25 30. A nucleotide in accordance with item 29 wherein said protein is a lectin.

30 31. A nucleotide in accordance with item 30 wherein said lectin is Concanavalin A.

32. A nucleotide in accordance with item 1 wherein said Sig chemical moiety comprises a-mannosyl residue and wherein said a-mannosyl residue is complexed with or bound to Concanavalin A.

05 33. A nucleotide in accordance with item 1 wherein said Sig chemical moiety comprises N-acetylglucosamine residue and wherein N-acetylglucosamine is complexed with or bound to wheat germ agglutinin.

10 34. A nucleotide in accordance with item 1 wherein said Sig chemical moiety includes an electron dense component.

15 35. A nucleotide in accordance with item 30 wherein said lectin comprises ferritin attached thereto.

36. A nucleotide in accordance with item 31 wherein said Concanavalin A is conjugated to ferritin.

20 37. A nucleotide in accordance with item 1 wherein said Sig chemical moiety includes or comprises a radioactive isotope.

25 38. A nucleotide in accordance with item 37 wherein said radioactive isotope is radioactive cobalt.

39. A nucleotide in accordance with item 1 wherein said Sig chemical moiety includes or comprises an enzyme.

30 40. A nucleotide in accordance with item 39 wherein said enzyme is alkaline phosphatase.

41. A nucleotide in accordance with item 39 wherein
said enzyme is acid phosphatase.

42. A nucleotide in accordance with item 1 wherein
05 said Sig chemical moiety includes or comprises a
fluorescing component attached thereto.

43. A nucleotide in accordance with item 42 wherein
said fluorescing component is fluorescein.

10

44. A nucleotide in accordance with item 1 wherein
said Sig chemical moiety includes or comprises a
magnetic component associated or attached thereto.

15 45. A nucleotide in accordance with item 44 wherein
said magnetic component comprises a magnetic oxide.

46. A nucleotide in accordance with item 45 wherein
said magnetic oxide is ferric oxide.

20

47. A nucleotide in accordance with item 1 wherein
said Sig chemical moiety includes an antigenic or hapten
component capable of complexing with an antibody
specific to said component.

25

48. A single-stranded polynucleotide comprising one or
more nucleotides in accordance with item 1.

49. A single-stranded polynucleotide in accordance with
30 item 48 wherein said single-stranded polynucleotide is
a polydeoxyribonucleotide.

50. A single-stranded polynucleotide in accordance with
it m 48 wherein said single-stranded polynucleotide is
35 a polyribonucleotide.

51. A double-stranded polynucleotide comprising one or more nucleotides in accordance with item 1.

52. A double-stranded polynucleotide in accordance with
05 item 51 wherein said double-stranded polynucleotide is
a double-stranded deoxyribonucleic acid.

53. A double-stranded polynucleotide in accordance with
item 51 wherein said double-stranded polynucleotide is
10 a double-stranded ribonucleic acid.

54. A double-stranded polynucleotide in accordance with
item 51 wherein said double-stranded polynucleotide is
a double-stranded deoxyribonucleic acid-ribonucleic acid
15 hybrid.

55. A polynucleotide comprising one or more nucleotides
in accordance with item 1 coupled or attached to a
polypeptide, said polypeptide having attached thereto
20 one or more biotin groups.

56. A polynucleotide in accordance with item 55
wherein said polypeptide is a polylysine.

25 57. A polynucleotide coupled or attached to a poly-
peptide, said polypeptide having attached thereto one or
more streptavidin units.

58. A polynucleotide terminally ligated or attached on
30 at least one end with a polypeptide, said polypeptide
having attached thereto one or more enzyme groups.

59. A single stranded polydeoxyribonucleotide coupled
or attached to a polypeptide, said polypeptide having
35 attached thereto one or more biotin groups.

60. A single-stranded polyribonucleotide coupled or attached to a polypeptide, said polypeptide having attached thereto one or more biotin groups.

05 61. A single-stranded polynucleotide comprising at least 12 nucleotides; at least one of said nucleotides being a nucleotide in accordance with item 1.

10 62. A polynucleotide coupled or attached to a polysaccharide.

63. A polynucleotide in accordance with item 62 wherein said polysaccharide has attached thereto or complexed therewith a plant binding protein.

15 64. A polynucleotide in accordance with item 63 wherein said protein is Concanavalin A.

65. A nucleotide in accordance with item 1 wherein
20 said base B is cytosine.

66. A nucleotide in accordance with item 1 wherein
said base B is uracil.

25 67. A nucleotide in accordance with item 1 wherein
said base B is thymine.

68. A nucleotide in accordance with item 1 wherein
said base B is adenine.

30 69. A nucleotide in accordance with item 1 wherein
said base B is guanine.

70. A nucleotide in accordance with item 1 wherein
35 said base B is 2-methyladenine.

-18-

71. A nucleotide in accordance with item 1 wherein
said base B is 1-methylguanine.

72. A nucleotide in accordance with item 1 wherein
05 said base B is 5-methylcytosine.

73. A nucleotide in accordance with item 1 wherein
said base B is 5-hydroxymethylcytosine.

10 74. A nucleotide in accordance with item 1 wherein
said Sig chemical moiety comprises a chelating agent.

75. A nucleotide in accordance with item 1 wherein
said base B is deazaadenine.

15 76. A nucleotide in accordance with item 1 wherein
said base B is deazaguanine.

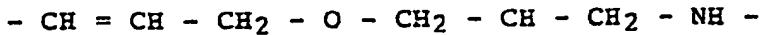
77. A nucleotide in accordance with item 1 wherein
20 said Sig chemical moiety
is connected to said base B via a chemical
linkage.

78. A nucleotide in accordance with item 77 wherein
25 said chemical linkage includes an olefinic bond at the
a-position relative to base B.

79. A nucleotide in accordance with item 77 wherein
said chemical linkage includes the moiety,
- CH₂ - NH - .

30 80. A nucleotide in accordance with item 77 wherein
said chemical linkage is,
- CH = CH - CH₂ - NH - .

81. A nucleotide in accordance with item 1 wherein the chemical moiety is,



05

OH .

82. A nucleotide in accordance with item 1 wherein said chemical linkage is selected from or includes a moiety selected from the group consisting of

10

O

"



15 83. A nucleotide in accordance with item 1 wherein said Sig chemical moiety includes or comprises a catalytic metal component.

84. A nucleotide in accordance with item 39 wherein
20 said enzyme is β -galactosidase.

85. A nucleotide in accordance with item 39 wherein said enzyme is glucose oxidase.

25 86. A nucleotide in accordance with item 39 wherein said enzyme is horseradish peroxidase.

87. A nucleotide in accordance with item 42 wherein said fluorescing component is fluorescein.

30

88. A nucleotide in accordance with item 42 wherein said fluorescing component is rhodamine.

89. A nucleotide in accordance with item 42 wherein
35 said fluorescing component is dansyl.

-20-

90. A nucleotide in accordance with item 80 wherein
said Sig chemical moiety is attached to the - NH - group
of said chemical linkage.

05 91. A nucleotide in accordance with item 90 wherein
said Sig chemical moiety comprises a polysaccharide.

92. A nucleotide in accordance with item 90 wherein
said Sig chemical moiety is biotin.

10 93. A nucleotide in accordance with item 90 wherein
said Sig chemical moiety is streptavidin.

15 94. A nucleotide in accordance with item 82 wherein
said Sig chemical moiety attached to said chemical
linkage is a monosaccharide.

20 95. A nucleotide in accordance with item 82 wherein
said Sig chemical moiety attached to said chemical
linkage is a streptavidin.

96. A polynucleotide in accordance with item 63
wherein said protein is a lectin.

25 97. A polynucleotide in accordance with item 62
wherein said polynucleotide is terminally ligated to
said polysaccharide.

30 98. A polyribonucleotide in accordance with item 60
wherein said polyribonucleotide is terminally ligated or
attached to said polypeptide.

35 99. A polydeoxyribonucleotide in accordance with item
59 wherein said polydeoxyribonucleotide is terminally
ligated or attached to said polypeptide.

100. A polynucleotide in accordance with item 55 wherein said polynucleotide is terminally ligated or attached to said polypeptide.

05 101. A ribonucleotide having the general formula,

Sig

P - S - B

wherein P is the phosphoric acid moiety, S the sugar
10 moiety and B the base moiety, the phosphoric acid moiety being attached at the 2', 3' and/or 5' position of the sugar moiety, said base B being attached from the N1 position or the N9 position to the 1' position of the sugar moiety when said base is a pyrimidine or a purine,
15 respectively, and wherein said Sig is a chemical moiety covalently attached to the sugar S, said Sig, when attached to said sugar S, being capable of signalling itself or making itself self-detecting or its presence known.

20

102. A ribonucleotide in accordance with item 101 wherein said Sig chemical moiety is attached to the C2' position of said sugar moiety.

25

103. A ribonucleotide in accordance with item 101 wherein said Sig chemical moiety is attached to the C3' position of said sugar moiety.

30

104. A ribonucleotide in accordance with item 101 wherein said Sig chemical moiety is attached to the S sugar moiety such that an oligoribonucleotide or poly-ribonucleotide containing said ribonucleotide is capable of forming a double-stranded ribonucleic acid or a DNA-RNA hybrid when said ribonucleotide is incorporated 35 into said oligoribonucleotide or said polyribonucleotide.

-22-

105. A polyribonucleotide comprising at least one ribonucleotide in accordance with item 101.

106. A ribonucleotide in accordance with item 1 wherein said base B is a pyrimidine.

05

107. A ribonucleotide in accordance with item 101 wherein said base B is a purine.

108. A ribonucleotide in accordance with item 101
10 wherein said base B is uracil.

109. A ribonucleotide in accordance with item 101
wherein said base B is adenine.

15 110. A ribonucleotide in accordance with item 101
wherein said base B is guanine.

111. A ribonucleotide in accordance with item 101
wherein said base B is cytosine.

20

112. A ribonucleotide in accordance with item 101
wherein said base B is a 7-deazapurine.

113. A ribonucleotide in accordance with item 101
25 wherein said Sig chemical moiety is a polysaccharide or
an oligosaccharide.

114. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety is a monosaccharide.

30

115. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety is a monosaccharide
selected from the group consisting of triose, tetrose,
pentose, hexose, heptose and octose.

116. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety is a sugar residue and
wherein said sugar residue is complexed with or attached
to a sugar or polysaccharide binding protein.

05

117. A ribonucleotide in accordance with item 116
wherein said protein is a lectin.

118. A ribonucleotide in accordance with item 117
10 wherein said lectin is Concanavalin A.

119. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety includes an electron
dense component.

15

120. A ribonucleotide in accordance with item 119
wherein said electron dense component is ferritin.

121. A ribonucleotide in accordance with item 101
20 wherein said Sig chemical moiety includes or comprises a
radioactive isotope component.

122. A ribonucleotide in accordance with item 121
wherein said radioactive isotope component is
25 radioactive cobalt.

123. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety comprises a catalytic
metal component.

30

124. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety includes or comprises
an enzyme.

125. A ribonucleotide in accordance with item 124
wherein said enzyme is alkaline phosphatase.

05 126. A ribonucleotide in accordance with item 124
wherein said enzyme is β -galactosidase.

10 127. A ribonucleotide in accordance with item 124
wherein said enzyme is glucose oxidase.

128. A ribonucleotide in accordance with item 124
wherein said enzyme is horseradish peroxidase.

15 129. A ribonucleotide in accordance with item 124
wherein said enzyme is ribonuclease.

130. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety includes or comprises a
floresching component attached thereto.

20 131. A ribonucleotide in accordance with item 130
wherein said fluorescing component is fluoroscein.

132. A ribonucleotide in accordance with item 130
wherein said fluorescing component is rhodamine.

25 133. A ribonucleotide in accordance with item 130
wherein said fluorescing component is dansyl.

30 134. A ribonucleotide in accordance with item 101
wherein said Sig chemical moiety comprises a magnetic
component associated or attached thereto.

135. A ribonucleotide in accordance with item 134
wherein said magnetic component comprises a magnetic
oxide.

136. A ribonucleotide in accordance with item 101 wherein said Sig chemical moiety includes an antigenic or hapten component capable of complexing with an antibody specific to said component.

05

137. A single stranded polyribonucleotide comprising one or more ribonucleotides in accordance with item 101, said single stranded polyribonucleotide comprising at least three said ribonucleotides.

10

138. A single stranded polyribonucleotide comprising at least twelve ribonucleotides and containing at least one ribonucleotide in accordance with item 101.

15

139. A polyribonucleotide coupled or attached to a polypeptide.

140. A polyribonucleotide in accordance with item 139 wherein said polypeptide is terminally attached or
20 ligated to said polyribonucleotide.

20

-26-

141. A nucleotide having the general formula

Sig



wherein P is the phosphoric acid moiety, S the sugar
05 moiety and B the base moiety, the phosphoric acid moiety
being attached to the 3' and/or the 5' position of the
sugar moiety when said nucleotide is a deoxyribonucleotide
and at the 2', 3' and/or 5' position when said nucleotide
10 is a ribonucleotide, said base B being a purine or
pyrimidine, said base B being attached from the N1
position or the N9 position to the 1' position of the
sugar moiety when said base B is a pyrimidine or a purine,
respectively, and wherein Sig is a chemical moiety
15 covalently attached to the phosphoric acid moiety via the
chemical linkage

OH

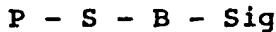


"

O

20 said Sig, when attached to said phosphoric acid moiety P
being capable of signalling itself or making itself self-
detecting or its presence known.

142. A nucleotide having the general formula



wherein P is the phosphoric acid moiety, S the sugar and monosaccharide moiety, B being the base moiety, the phosphoric acid moiety being attached to the 3' and/or

05 the 5' position of the sugar moiety when said nucleotide is deoxyribonucleotide and at the 2', 3' and/or 5' position when said nucleotide is a ribonucleotide, said base being a purine or a pyrimidine, said base being attached from the N1 position or the N9 position to the
10 Cl' position of the sugar moiety when said base is a pyrimidine or a purine, respectively, and wherein said Sig is a chemical moiety covalently attached to the base B of said nucleotide, said Sig being attached to the N⁶ or 6-amino group when said base B is adenine or the N² or 2-amino group when said base B is guanine or the N⁴ or 4-amino group when said base B is cytosine, said Sig when attached to said base B being capable of signally itself or makes itself self-detecting or its presence known.

143. A nucleotide having the general formula P-S-B,
wherein P is the phosphoric acid moiety, S the sugar or
monosaccharide moiety and B the base moiety, said
nucleotide having covalently attached to the P or S or B
moiety a chemical moiety Sig, said Sig chemical moiety
when attached to the P moiety is attached thereto via the
chemical linkage,

05

10

OH

"

- P - O - Sig,

"

O

and when Sig is attached to the S moiety, the S moiety is
a ribose group, said chemical moiety Sig when attached to
15 said P, S or B being capable of signalling itself or makes
itself self-detecting or its presence known.

20

144. A nucleotide in accordance with item 143 wherein
said nucleotide is capable of being incorporated into to
form a double-stranded ribonucleic acid, a double-
stranded deoxyribonucleic acid or a double-stranded
deoxyribonucleic acid-ribonucleic acid hybrid.

25

145. A nucleotide in accordance with item 143 wherein
when said nucleotide is incorporated into or attached to a
double-stranded deoxyribonucleic acid or double- stranded
ribonucleic acid or DNA-RNA hybrid, said chemical moiety
Sig is capable of signalling itself or making itself
self-detecting or its presence known.

30

146. A single-stranded polynucleotide comprising one or
more nucleotides in accordance with item 143.

147. A double-stranded polynucleotide comprising one or
35 more nucleotides in accordance with 143.

148. A single-stranded polydeoxyribonucleotide containing at least 12 nucleotides and comprising one or more nucleotides in accordance with item 143.

05 149. A nucleotide in accordance with Claim 143 wherein said Sig moiety is attached to the B base moiety through a mono- or oligosaccharide linkage.

10 150. A nucleotide in accordance with item 143 wherein said Sig moiety is monosaccharide or polysaccharide moiety attached to said B base moiety.

15 151. A polynucleotide comprising at least one nucleotide in accordance with item 143.

152. A polynucleotide comprising at least one nucleotide in accordance with item 149.

20 153. A polynucleotide comprising at least one nucleotide in accordance with item 150.

154. A nucleotide in accordance with Claim 143 or polynucleotide comprising at least one nucleotide in accordance with item 143 wherein said Sig chemical moiety comprises an agent for stimulating or inducing the production of interferon.

30 155. A method of chemotherapy suitable for inhibiting RNA and/or DNA synthesis which comprises administering to an organism capable of and/or functioning for the production or synthesis of DNA and/or RNA an effective DNA and/or RNA synthesis inhibiting amount of a nucleotide in accordance with item 143.

156. A method of chemotherapy suitable for inhibiting RNA and/or DNA synthesis which comprises administering to an organism capable of and/or functioning for the production or synthesis of DNA and/or RNA an effective DNA and/or RNA synthesis inhibiting amount of a nucleotide in accordance
05 with item 1.

157. A method of chemotherapy suitable for inhibiting RNA and/or DNA synthesis which comprises administering to the organism capable of and/or functioning for the production or synthesis of DNA and/or RNA an effective DNA and/or RNA synthesis inhibiting amount of a nucleotide in accordance
10 with item 101.

158. A method of chemotherapy suitable for inhibiting RNA and/or DNA synthesis which comprises administering to the organism capable of and/or functioning for the production or synthesis of DNA and/or RNA an effective DNA and/or RNA synthesis inhibiting amount of a nucleotide in accordance
20 with item 141.

159. A method of chemotherapy suitable for inhibiting RNA and/or DNA synthesis which comprises administering to the organism capable of and/or functioning for the production or synthesis of DNA and/or RNA an effective DNA and/or RNA synthesis inhibiting amount of a nucleotide in accordance
25 with item 142.

160. A method of chemotherapy in accordance with item 155 wherein the B base moiety of said nucleotide is glycosylated.
30

161. A method of chemotherapy in accordance with item 155 wherein the Sig chemical moiety of said nucleotide comprises an anti-tumor or cytotoxic agent.

162. A method for the stimulation or induction of cells
for the production of lymphokines, cytokinins and/or
interferon which comprises introducing into or bringing
into contact with cells capable of and/or functioning for
05 the production of said lymphokines, cytokinins and/or
interferon an effective lymphokine, cytokinin and/or
interferon stimulating and production inducing amount of a
nucleotide in accordance with item 143.

10 163. A method for the stimulation or induction of cells
for the production of lymphokines, cytokinins and/or
interferon which comprises introducing into or bringing
into contact with cells capable of and/or functioning for
15 the production of said lymphokines, cytokinins and/or
interferon an effective lymphokine, cytokinin and/or
interferon stimulating and production inducing amount of a
nucleotide in accordance with item 1.

20 164. A method for the stimulation or induction of cells
for the production of lymphokines, cytokinins and/or
interferon which comprises introducing into or bringing
into contact with cells capable of and/or functioning for
the production of said lymphokines, cytokinins and/or
25 interferon an effective lymphokine, cytokinin and/or
interferon stimulating and production inducing amount of a
nucleotide in accordance with item 101.

165. A method for the stimulation or induction of cells
for the production of lymphokines, cytokinins and/or
interferon which comprises introducing into or bringing
into contact with cells capable of and/or functioning for
the production of said lymphokines, cytokinins and/or
interferon an effective lymphokine, cytokinin and/or
interferon stimulating and production inducing amount of a
nucleotide in accordance with item 141.

166. A method for the stimulation or induction of cells
for the production of lymphokines, cytokinins and/or
interferon which comprises introducing into or bringing
into contact with cells capable of and/or functioning for
the production of said lymphokines, cytokinins and/or
interferon an effective lymphokine, cytokinin and/or
interferon stimulating and production inducing amount of a
nucleotide in accordance with item 142.

167. A polynucleotide comprising one or more nucleotides
in accordance with item 1 or item 101 or item 141 or
item 142 or item 143, coupled or attached to a poly-
peptide, said polypeptide having attached thereto one or
more Sig chemical moieties, said Sig when attached to said
polypeptide being capable of signalling itself or making
itself self-detecting or its presence known.

168. A polynucleotide coupled or attached to a poly-
peptide, said polypeptide having attached thereto one or
more Sig chemical moieties, said Sig chemical moieties
when attached to said polypeptide being capable of
signalling itself or making itself self-detecting or
making its presence known.

-33-

169. A nucleotide in accordance with item 1 or item 101 or item 141 or item 142 or item 143 wherein said Sig chemical moiety includes a chelating agent.

05 170. A nucleotide in accordance with item 169 wherein said chelating agent includes the chemical moiety

10

15 wherein M is H or a substitutable metal.

171. A nucleotide in accordance with item 170 wherein said metal is magnesium or a metal replaceable by cobalt.

20 172. A polynucleotide containing one or more nucleotides in accordance with item 1 or item 101 or item 141 or item 142 or item 143 wherein said Sig chemical moiety includes a chelating agent.

25 173. A polynucleotide containing a nucleotide in accordance with item 1 or item 101 or item 141 or item 142 or item 143 wherein said Sig chemical moiety includes a chelating agent in accordance with Claim 170.

30 174. A nucleotide in accordance with item 170 wherein said metal is a catalytically active metal.

175. A nucleotide in accordance with item 170 wherein said metal is a heavy metal.

-34-

176. A nucleotide in accordance with item 170 wherein said metal is radioactive cobalt.

177. A nucleotide in accordance with item 170 wherein
05 said metal is magnesium.

178. A ribonucleotide in accordance with item 1 or item 101 or item 141 or item 142 or item 143 wherein said Sig chemical moiety includes a chelating agent.

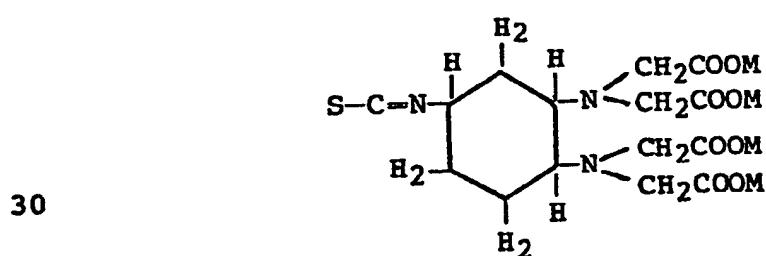
10 179. A deoxyribonucleotide in accordance with item 1 or item 101 item 141 or item 142 or item 143 wherein said Sig chemical moiety includes a chelating agent.

15 180. A nucleotide in accordance with item 170 wherein said metal M is a radioactive isotope.

181. A nucleotide in accordance with item 170 wherein said metal is platinum.

20 182. A nucleotide in accordance with item 170 wherein said M is hydrogen or a substitutable metal or a radioactive element.

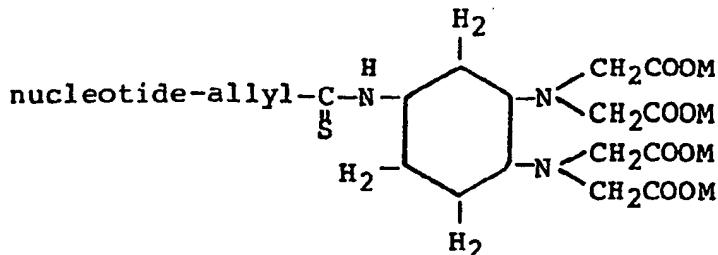
25 183. The compound



wherein M is hydrogen or a metal.

184. The compound

05



wherein M is hydrogen or a metal.

10

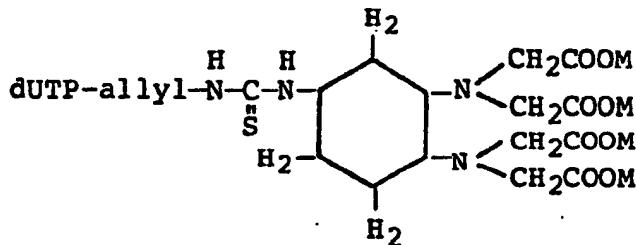
185. A compound in accordance with item 184 wherein said nucleotide is a deoxyribonucleotide.

15

186. A compound in accordance with item 184 wherein said nucleotide is a ribonucleotide.

187. The compound

20



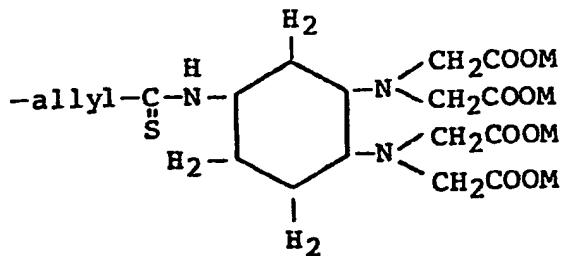
25

wherein M is hydrogen or a metal.

-36-

188. A nucleotide in accordance with items
1, 101, 141, 142 or 143, wherein the Sig
chemical moiety comprises the chemical moiety

05



10

189. A polynucleotide containing one or more nucleotides
in accordance with item 188.

190. A polynucleotide in accordance with item 189
15 wherein said polynucleotide is a single-stranded deoxy-
ribonucleotide or a single-stranded ribonucleotide or a
double-stranded DNA, RNA or DNA-RNA hybrid.

20

191. A nucleotide in accordance with item 143 wherein
said Sig moiety comprises a saccharide component, a
protein component capable of binding to said saccharide
component and a glycosylated enzyme component capable of
binding to said protein component.

05

192. A nucleotide in accordance with item 191 wherein
said saccharide component is a monosaccharide.

10 193. A nucleotide in accordance with item 191 wherein
said saccharide component is an oligosaccharide.

194. A nucleotide in accordance with item 191 wherein
said saccharide component is a polysaccharide.

15 195. A nucleotide in accordance with item 191 wherein
said protein component is a lectin.

20 196. A nucleotide in accordance with item 191 wherein
said lectin is a plant lectin selected from the group
consisting of Concanavalin A, LCH lentil lectin, PSA pea
lectin and BFA Vicia Faba lectin.

25 197. A nucleotide in accordance with item 191 wherein
said enzyme component is an enzyme selected from the group
consisting of alkaline phosphatase, acid phosphatase and
horseradish peroxidase.

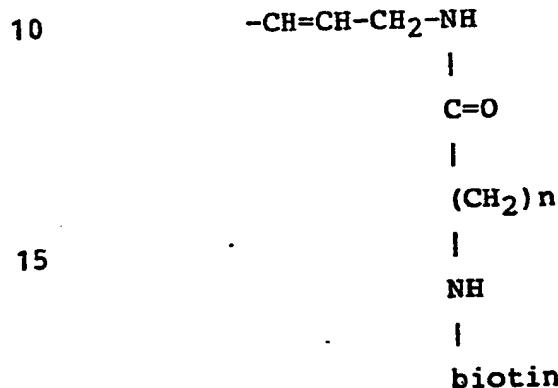
30 198. An amino acid or polypeptide comprising a Sig moiety
attached thereto.

199. An amino acid or polypeptide in accordance with
item 198 wherein said Sig moiety comprises a saccharide
component.

200. A monosaccharide or a polysaccharide comprising a
Sig moiety attached thereto.

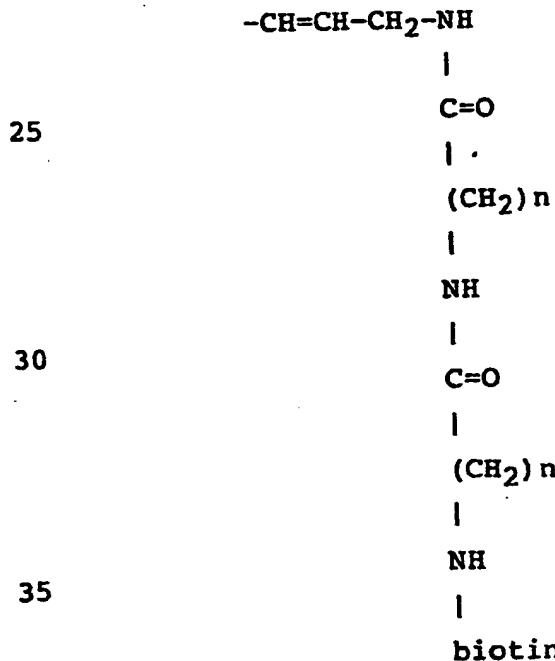
201. A monosaccharide or a polysaccharide in accordance
05 with item 200 wherein said Sig moiety comprises a
chelating agent.

202. A nucleotide in accordance with item 143 wherein
said Sig moiety comprises the linkage



wherein n is an integer having a value in the range 2-10.

20 203. A nucleotide in accordance with item 143 wherein
said Sig moiety comprises the linkage



wherein n is an integer having a value in the range 2-10.

--- 204. A method of detecting a first compound which includes a nucleotide in accordance with item 1 as part of said first compound, which comprises contacting said first compound with a second compound capable of forming a complex therewith under suitable conditions so as to form said complex, said complex comprising said first compound and said second compound and detecting said complex. --

--- 205. A method in accordance with item 204 wherein said second compound is a polynucleotide. --

-- 206. A method of detecting a first compound which includes a ribonucleotide in accordance with item 101 as part of said first compound, which comprises contacting said first compound with a second compound capable of forming a complex therewith under conditions so as to form said complex and detecting said complex. --

-- 207. A method of detecting a first compound which includes a nucleotide in accordance with item 141 as part of said first compound which comprises contacting said first compound with a second compound capable of forming a complex therewith under suitable conditions so as to form said complex and detecting said complex. --

-- 208. A method of detecting a first compound which includes a nucleotide in accordance with item 142 as part of said first compound which comprises contacting said first compound with a second compound capable of forming a complex therewith under suitable conditions so as to form said complex and contacting said complex. --

-- 209. A method of detecting a first compound which includes a nucleotide in accordance with item 143 as part of said first compound which comprises contacting said first compound with a second compound capable of forming a complex therewith under suitable conditions so as to form said complex and detecting said complex. --

-- 210. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a nucleotide in accordance with item 1 and detecting said duplex. --

-- 211. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a ribonucleotide in accordance with item 101 and detecting said duplex. --

-- 212. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a nucleotide in accordance with item 141 and detecting said duplex. --

-- 213. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a

double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a nucleotide in accordance with item 142 and detecting said duplex. --

-- 214. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a nucleotide in accordance with item 143 and detecting said duplex. --

-- 215. A method of detecting the presence of a nucleic acid-containing etiological agent in a subject which comprises obtaining a suitable sample from said subject, detecting the presence in said sample of a deoxyribonucleic or ribonucleic acid naturally associated with said etiological agent by forming under suitable conditions a double-stranded polynucleotide duplex which includes a compound in accordance with item 1 and a single strand of deoxyribonucleic or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid which is naturally associated with said etiological agent under suitable conditions and detecting the presence of said double-stranded polynucleotide duplex. --

-- 216. A method of detecting the presence of a nucleic acid-containing etiological agent in a subject which comprises obtaining a suitable sample from said subject, detecting the presence in said sample of a deoxyribonucleic or ribonucleic acid naturally associated with said etiological agent by forming under suitable conditions a double-stranded polynucleotide duplex which includes a compound in accordance with item 101 and a single strand of deoxyribonucleic or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid which is naturally associated with said etiological agent under suitable conditions and detecting the presence of said double-stranded polynucleotide duplex. --

-- 217. A method of detecting the presence of a nucleic acid-containing etiological agent in a subject which comprises obtaining a suitable sample from said subject, detecting the presence in said sample of a deoxyribonucleic or ribonucleic acid naturally associated with said etiological agent by forming under suitable conditions a double-stranded polynucleotide duplex which includes a compound in accordance with item 141 and a single strand of deoxyribonucleic or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid which is naturally associated with said etiological agent under suitable conditions and detecting the presence of said double-strand d polynucleotide duplex. -

-- 218. A method of detecting the presence of a nucleic acid-containing etiological agent in a subject which comprises obtaining a suitable sample from said subject, detecting the

presence in said sample of a deoxyribonucleic or ribonucleic acid naturally associated with said etiological agent by forming under suitable conditions a double-stranded polynucleotide duplex which includes a compound in accordance with item 142 and a single strand of deoxyribonucleic or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid which is naturally associated with said etiological agent under suitable conditions and detecting the presence of said double-stranded polynucleotide duplex. --

-- 219. A method of detecting the presence of a nucleic acid-containing etiological agent in a subject which comprises obtaining a suitable sample from said subject, detecting the presence in said sample of a deoxyribonucleic or ribonucleic acid naturally associated with said etiological agent by forming under suitable conditions a double-stranded polynucleotide duplex which includes a compound in accordance with item 143 and a single strand of deoxyribonucleic or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid which is naturally associated with said etiological agent under suitable conditions and detecting the presence of said double-stranded polynucleotide duplex. --

-- 220. A method in accordance with item 215 wherein said nucleic acid-containing etiological agent is a bacterium, a virus or a fungus. --

-- 221. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with item 1 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

-- 222.. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with item 101 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

-- 223. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with item 141 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

-- 224. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with item 142 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

--225. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with item 143 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

-- 226. A method in accordance with items 221, 222, 223, 224 or 225 wherein said bacterium is selected from the group consisting of Streptococcus pyogenes and Neisseria meningitidis and said antibiotic a penicillin. --

-- 227. A method in accordance with item 221 wherein said bacterium is selected from the group consisting of Staphylococcus aureus, Candida albicans, Pseudomonas aeruginosa, Streptococcus pyogenes and Neisseria gonorrhoeae and said antibiotic a tetracycline. --

-- 228. A method in accordance with item 221 wherein said bacterium is Mycobacterium tuberculosis and said antibiotic is an amino glycoside. --

-- 229. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said subject which is associated with said genetic disorder and includes a compound in accordance with item 1 incorporated therein, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid duplex indicating the presence or absence of said genetic disorder. --

-- 230. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said subject which is associated with said genetic disorder and includes a compound in accordance with item 101 incorporated therein, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid duplex indicating the presence or absence of said genetic disorder. --

-- 231. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said subject which is associated with said genetic disorder and includes a

compound in accordance with item 141 incorporated therein, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid duplex indicating the presence or absence of said genetic disorder. --

-- 232. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said subject which is associated with said genetic disorder and includes a compound in accordance with item 142 incorporated therein, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid duplex indicating the presence or absence of said genetic disorder. --

-- 233. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleotide acid gene sequence of said subject which is associated with said genetic disorder and includes a compound in accordance with item 143 incorporated therein, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid

duplex indicating the presence or absence of said genetic disorder. --

-- 234. A method of diagnosing thalassemia in a human subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence which is absent in β -minus-thalassemia subjects and includes a nucleotide in accordance with item 1, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and determining the presence of said duplex, the absence of said duplex indicating the presence of β -minus-thalassemia. --

-- 235. A method of diagnosing thalassemia in a human subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence which is absent in β -minus-thalassemia subjects and includes a nucleotide in accordance with item 101, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and determining the presence of said duplex, the absence of said duplex indicating the presence of β -minus-thalassemia. --

-- 236. A method of diagnosing thalassemia in a human subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence which is absent in β -minus-thalassemia subjects and includes a nucleotide in accordance with item 141, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from

| said subject so as to form a double-stranded hybrid duplex and
determining the presence of said duplex, the absence of said
duplex indicating the presence of β -minus-thalassemia. -- -

-- 237. A method of diagnosing thalassemia in a human subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence which is absent in β -minus-thalassemia subjects and includes a nucleotide in accordance with item 142, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and determining the presence of said duplex, the absence of said duplex indicating the presence of β -minus-thalassemia. --

-- 238. A method of diagnosing thalassemia in a human subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence which is absent in β -minus-thalassemia subjects and includes a nucleotide in accordance with item 143, contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and determining the presence of said duplex, the absence of said duplex indicating the presence of β -minus-thalassemia. --

-- 239. A method of chromosomal karyotyping which comprises preparing a series of modified polynucleotides corresponding to a series of defined genetic sequences located on chromosomes, said polynucleotides including one or more compounds in accordance with item 1, contacting said polynucleotides with deoxyribo-

nucleic acid of or obtained from chromosomes so as to form hybrid duplexes and detecting said duplexes, thereby determining the location of said duplexes on said chromosomes and the location of said genetic sequences on said chromosomes. --

-- 240. A method of chromosomal karyotyping which comprises preparing a series of modified polynucleotides corresponding to a series of defined genetic sequences located on chromosomes, said polynucleotides including one or more compounds in accordance with item 101, contacting said polynucleotides with deoxyribonucleic acid of or obtained from chromosomes so as to form hybrid duplexes and detecting said duplexes, thereby determining the location of said duplexes on said chromosomes and the location of said genetic sequences on said chromosomes. --

-- 241. A method of chromosomal karyotyping which comprises preparing a series of modified polynucleotides corresponding to a series of defined genetic sequences located on chromosomes, said polynucleotides including one or more compounds in accordance with item 141, contacting said polynucleotides with deoxyribonucleic acid of or obtained from chromosomes so as to form hybrid duplexes and detecting said duplexes, thereby determining the location of said duplexes on said chromosomes and the location of said genetic sequences on said chromosomes. --

-- 242. A method of chromosomal karyotyping which comprises preparing a series of modified polynucleotides corresponding to a series of defined genetic sequences located on chromosomes, said polynucleotides including one or more compounds in accordance with item 142, contacting said polynucleotides with deoxyribonucleic acid of or obtained from chromosomes so as to form hybrid duplexes and detecting said duplexes, thereby determining the location of said duplexes on said chromosomes and the location of said genetic sequences on said chromosomes. --

-- 243. A method of chromosomal karyotyping which comprises preparing a series of modified polynucleotides corresponding to a series of defined genetic sequences located on chromosomes, said polynucleotides including one or more compounds in accordance with item 143, contacting said polynucleotides with deoxyribonucleic acid of or obtained from chromosomes so as to form hybrid duplexes and detecting said duplexes, thereby determining the location of said duplexes on said chromosomes and the location of said genetic sequences on said chromosomes. --

-- 244. A method of identifying or locating hormone receptor sites on the surface of cells which comprises binding a compound in accordance with items 1, 101, 141, 142 or 143
| to said sites under suitable conditions
| permitting binding of said compound to said receptor sites and
detecting said compounds bound to said receptor sites. --

-54-

-- 245. A method of tumor or cancer cell identification or detection which comprises identifying or detecting malignant cells by detecting abnormal receptor sites associated therewith in accordance with item 244. --

-- 246. A method of diagnosing a tumor cell which comprises preparing a polynucleotide which is complementary to a messenger ribonucleic acid synthesized from a deoxyribonucleic acid gene sequence associated with the production of a poly-peptide diagnostic for or identifiable with said tumor cell or which is complementary to said deoxyribonucleic acid gene sequence and which includes a compound in accordance with items 1, 101, 141, 142 or 143,

introducing said polynucleotide into contact with said cell under suitable conditions so as to permit said polynucleotide to hybridize with said deoxyribonucleic acid gene sequence or said messenger ribonucleic acid and detecting the formation of a resulting formed hybrid containing said polynucleotide. --

-- 247. A diagnostic kit useful for determining the presence of a nucleic acid-containing organism or the like which comprises a polynucleotide which includes in its make-up a compound selected from the compounds of items 1, 101, 141, 142 or 143, and which is complementary to all or an identifiable, distinct or unique portion of the nucleic acid contained in said organism and means for detecting or expressing the presence or absence of a resulting formed hybrid between said polynucleotide and said nucleic acid of said organism

when said polynucleotide is brought into contact with the nucleic acid of said organism or the like under hybrid forming conditions. --

-- 248. A method of determining or identifying or diagnosing a first compound capable of complexing with or binding with a first component of a second compound, said second compound comprising said first component and a second component, said second component being attached to or complexed with said first component, which comprises bringing said first compound into contact with said second compound whereby said first component of said second compound complexes with said first compound to form a third compound comprising said first compound and said second compound and contacting said resulting third compound with a fourth compound capable of binding with or attaching to said second component of said second compound making up said third compound. --

-- 249. A method in accordance with item 248 wherein multiple amounts of said second compound are separately brought into contact with said first compound or vice versa. --

-- 250. A method in accordance with item 248 wherein said first compound is a molecule with a receptor site that specifically binds to a second molecule, wherein said first component of said second compound is capable of specifically binding with said receptor site of said first compound, wherein said second component of said second compound is a sugar, oligosaccharide or polysaccharide and wherein said fourth compound is a polypeptide. --

-- 251. A method of determining or identifying or diagnosing a first compound capable of complexing with or binding with an amino acid, peptide or protein which comprises bringing said first compound into contact with a second compound, said second compound comprising an amino acid, peptide or protein attached to or complexed with a sugar, or oligosaccharide or polysaccharide, whereupon said second compound complexes with or becomes attached to said first compound to form a third compound comprising said first compound and said second compound and contacting said resulting third compound with a protein or amino acid or peptide capable of binding with or attaching to the sugar or oligosaccharide moiety of said second compound making up said third compound. --

-- 252. A method of determining or identifying or diagnosing a first compound capable of complexing or binding with a first component of a second compound, said second compound comprising said first component, a second component attached to or complexed with said first component, said second compound being a chelating agent, and a third component, said third component being an ion chelated or fixed to said second component, which comprises bringing said first compound into contact with said second compound whereby said first component of said compound complexes with said first compound to form a third compound comprising said first compound and said second compound. --

-- 253. A method in accordance with item 252 wherein said third compound is brought into contact with or reacted with a fourth compound under conditions such that said third component of said second compound making up said third compound catalyzes a discernible or detectable reaction involving or otherwise interacting with said fourth compound. --

-- 254. A method of chemotherapy involving the delivery of a therapeutic or cytotoxic agent to a selected receptor site of an organism or human subject which comprises bringing a first compound comprising a first component, a second component and a third component into contact with said receptor site, said first component being capable of attaching to or complexing with said receptor site, said second component being attached to said first component and said third component comprising a therapeutic or cytotoxic agent attached to said second component whereby upon contacting said receptor site with said first compound, said therapeutic or cytotoxic agent brought into contact with or delivered to said receptor site, said second component being a sugar, oligosaccharide or polypeptide and said third component containing said therapeutic or cytotoxic agent additionally comprising an amino acid or peptide or protein attached to or fixed to said second component. --

-- 255. A method of determining or identifying or diagnosing in an organism, cellular material or tissue a first compound comprising a receptor molecule with a receptor site that specifically binds to a second molecule which comprises bringing said organism, material or tissue containing said receptor molecule into contact with a second compound, said second compound comprising a first component capable of complexing with or binding with said receptor molecule, said second compound also comprising a second component, said second component being attached to or complexed with said first component, said first component of said second compound when in contact with said receptor molecule complexes with said receptor molecule to form a third compound comprising said first compound and said second compound and contacting said third compound with a fourth compound capable of binding with or attaching to said second compound making up said third compound, component of said second compound being selected said first component of said second compound being selected from group A consisting of an amino acid, a peptide, or a protein or from group B consisting of a sugar, oligosaccharide or polysaccharide or from group C consisting of a purine, or pyrimidine, nucleoside, nucleotide, oligonucleotide or polynucleotide, said second component of said second compound being a sugar, oligosaccharide or polysaccharide and said fourth compound is selected from the group consisting of a peptide or polypeptide, a lectin, or an antibody. --

-- 256. A nucleotide in accordance with item 1 wherein said Sig chemical moiety covalently attached to the base B of said nucleotide includes a coenzyme. --

-- 257. A nucleotide in accordance with item 256 wherein said coenzyme is selected from the group consisting of thiamine pyrophosphate, flavine mononucleotide, flavine adenine dinucleotide, nicotinamide adenine dinucleotide, nicotinamide adenine dinucleotide phosphate, coenzyme A, pyridoxyl phosphate, biotin, tetrahydrofolic acid, coenzyme B₁₂, lipoic acid and ascorbic acid. --

-- 258. A method of detecting a first compound which includes in its make-up a nucleotide in accordance with item 256 which comprises contacting said first compound with an apoenzyme corresponding to said coenzyme. --

-- 259. A method in accordance with item 258 wherein said coenzyme is flavine adenine dinucleotide (FAD) and wherein said apoenzyme is flavine adenine dinucleotide reductase. --

-- 260. A ribonucleotide in accordance with item 101 wherein said Sig chemical moiety includes a coenzyme. --

-- 261. A nucleotide in accordance with item 141 wherein said Sig chemical moiety includes a coenzyme. --

-- 262. A nucleotide in accordance with item 142 wherein said Sig chemical moiety includes a coenzyme. --

-- 263. A nucleotide in accordance with item 143 wherein said Sig chemical moiety includes a coenzyme. --

-- 264. A method of detecting a first compound which includes in its make-up a nucleotide in accordance with item 263 which comprises contacting said first compound with an apoenzyme corresponding to said coenzyme. --

--265. An amino acid or polypeptide in accordance with item 198, wherein said Sig moiety attached to said amino acid or polypeptide includes a coenzyme.--

--266. An amino acid or polypeptide in accordance with item 265, wherein said coenzyme is selected from the group consisting of thiamine pyrophosphate, flavine mononucleotide, flavine adenine dinucleotide, nicotinamide adenine dinucleotide, nicotinamide adenine dinucleotide phosphate, coenzyme A, pyridoxyl phosphate, biotin, tetrahydrofolic acid, coenzyme B₁₂, lipoic acid and ascorbic acid.--

--267. A method of detecting a first compound which includes in its make-up an amino acid or polypeptide in accordance with item 265, which comprises contacting said first compound with an apoenzyme corresponding to said coenzyme.--

--268. A method in accordance with item 267, wherein said coenzyme is flavin adenine dinucleotide (FAD) and wherein said apoenzyme is flavine adenine dinucleotide reductase.--

--269. A monosaccharide or polysaccharide in accordance with item 200, wherein said Sig moiety attached to said monosaccharide or said polysaccharide includes a coenzyme.--

--270. A monosaccharide or polysaccharide in accordance with item 269, wherein said coenzyme is selected from a group consisting of thiamine pyrophosphate, flavine mononucleotide,

flavine adenine dinucleotide, nicotinamide adenine dinucleotide, nicotinamide adenine dinucleotide phosphate, coenzyme A, pyridoxyl phosphate, biotin, tetrahydrofolic acid, coenzyme B₁₂, lipoic acid and ascorbic acid.--

271. A method of detecting a first compound which includes in its make-up a monosaccharide or polysaccharide in accordance with item 269, which comprises contacting said first compound with an apoenzyme corresponding to said coenzyme.--

272. A method in accordance with item 271, wherein said coenzyme is flavine adenine dinucleotide (FAD) and wherein said apoenzyme is flavine adenine dinucleotide reductase.--

-- 273. A nucleotide in accordance with item 1, wherein said Sig component comprises a radioactive component.

274. A nucleotide in accordance with item 273, wherein said radioactive component comprises radioactive labeled avidin attached thereto.

275. A nucleotide in accordance with item 273, wherein said Sig component comprises radioactive labeled streptavidin attached to biotin.

276. A single-stranded polynucleotide comprising one or more nucleotides in accordance with item 1, wherein said Sig component is radioactive.

277. A double-stranded polynucleotide comprising one or more nucleotides in accordance with item 1 and wherein said Sig component thereof is radioactive.

278. A nucleotide in accordance with item 101, wherein said Sig component comprises a radioactive moiety.

279. A polynucleotide comprising a ribonucleotide in accordance with item 101 wherein said Sig moiety is radioactively labeled.

280. A nucleotide in accordance with item 141, wherein said Sig moiety thereof is radioactively labeled.

281. A polynucleotide comprising one or more nucleotides in accordance with item 141 wherein said Sig moiety thereof is radioactively labeled.

282. A nucleotide in accordance with item 142, wherein said Sig moiety thereof is radioactively labeled.

283. A polynucleotide comprising one or more nucleotides in accordance with item 142 wherein said Sig moiety thereof is radioactively labeled.

284. A nucleotide in accordance with item 143, wherein said Sig moiety is radioactively labeled.

285. A polynucleotide comprising one or more nucleotides in accordance with item 143 wherein said Sig moiety thereof is radioactively labeled.

286. A single-stranded polynucleotide comprising one or more nucleotides in accordance with item 143 wherein said Sig component thereof is radioactively labeled.

287. A single-stranded polydeoxyribonucleotide in accordance with item 148, wherein said Sig component thereof is radioactively labeled.

288. A polynucleotide containing one or more nucleotides in accordance with item 1, or item 101, or item 141, or item 142, or item 143 wherein said Sig moiety is radioactive labeled.

289. A polynucleotide containing one or more nucleotides in accordance with item 1 or item 101, or item 141, or item 142, or item 143, wherein said Sig chemical moiety comprises biotin attached to radioactively labeled avidin.

290. A polynucleotide containing one or more nucleotides in accordance with item 1 or item 101, or item 141, or item 142, or item 143, wherein said Sig chemical moiety comprises biotin attached to radioactively labeled strepavidin.

291. A polynucleotide containing one or more nucleotides in accordance with item 1, or item 101, or item 141, or item 142, or item 143, wherein said Sig chemical moiety comprises biotin and radioactively labeled avidin or strepavidin attached to said biotin.

292. A method of determining the presence of a polynucleotide, comprising one or more nucleotides in accordance with item 1, or item 101, or item 141, or item 142, or item 143, wherein said Sig component of said nucleotide comprises biotin which comprises bringing said polynucleotide into contact with radioactive labeled avidin to bind said biotin to said radioactive labeled avidin and determining the presence of the biotin bound to said radioactive labeled avidin by detecting the radioactivity of the biotin-bound radioactive avidin.

293. A method of determining the presence of a polynucleotide, comprising one or more nucleotides in accordance with item 1, or item 101, or item 141, or item 142, or item 143, wherein said Sig component of said nucleotide comprises biotin which comprises bringing said polynucleotide

1 into contact with radioactive labeled strepavidin to bind
said biotin to said radioactive labeled strepavidin and
determining the presence of the biotin bound to said radio-
active labeled strepavidin by detecting the radioactivity
of the biotin-bound radioactive strepavidin.
5

294. A polynucleotide containing one or more nucleotides
in accordance with item 1 or item 101, or item 141, or
10 item 142, or item 143, wherein said Sig moiety comprises a
radioactively labeled antibody.

295. A polynucleotide containing one or more nucleotides
in accordance with item 1 or item 101, or item 141, or
15 item 142, or item 143, wherein said Sig moiety comprises
a radioactively labeled protein.

296. A polynucleotide containing one or more nucleotide:
20 in accordance with item 1 or item 101, or item 141, or
item 142, or item 143, wherein said Sign moiety comprises a
radioactively labeled lectin. --

25

30

35

The following examples are illustrative of various embodiments of the practices of this invention:

EXAMPLE I

05

Biotinyl-N-hydroxysuccinide ester (BNHS) was prepared according to a method of Becker et al, P.N.A.S. 68 2604 (1971). Biotin (0.24 g, 1.0 mmol) was dissolved in 5ml dry dimethylformamide. Dicyclohexylcarbodiimide (0.21 g, 10 1.0 mmol) and N-hydroxysuccinimide (.12 g, 1.0 mmol) were added and the solution stirred at room temperature for 15 hours. After filtration of the subsequent precipitate, the filtrate was evaporated at reduced pressure the residue was washed twice with ethanol and 15 recovered from hot isopropyl alcohol to yield a white crystalline product having a m.p. of 216-218°C.

EXAMPLE II

20 Biotinyl-1,6-diaminohexane amide was prepared as follows: A solution of 1,6-diaminohexane (320 mg, 2.0 mmol), dissolved in 50 ml water, was brought to pH 8.5 by addition of carbon dioxide. Biotinyl-N-hydroxy-succinimide ester (100 mg, 0.29 mmol), dissolved in 10 ml dimethylformamide, was added. After 18 hours at room 25 temperature the mixture was evaporated and the residue washed with ether and subsequently dried in a dessicator.

EXAMPLE III

Polybiotinylated poly-L-lysine was prepared by the
following procedure. Polylysine (100 umol lysine)
dissolved in 2 ml 0.1 M sodium borate, pH 8.5 was added
05 to biotinyl-N-hydroxysuccimide ester (17.5 mg, 50 umol)
dissolved in 0.5 ml dimethylformamide. After stirring
at room temperature for 18 hours, the mixture was
dialyzed against 10 mM tris buffer, pH 7.5.

10 EXAMPLE IV

Oligodeoxyribonucleotides were end-labeled using
cytidine-5'-triphosphate and terminal transferase as
follows. Purified phage DNA, alkali sheared with 0.2 N
15 sodium hydroxide and diluted to 2 A₂₆₀ units/ml in
potassium cacodylate (0.1 M), tris base (25 mm), cobalt
chloride (1 mM) and dithiothreitol (0.2 M) were used.
To this DNA solution (1 ml) was added cytidine-5'-tri-
phosphate (10 mmol) and terminal transferase (200 units).

20 After incubating at 37° for 5 to 8 hours the reaction
was stopped by the addition of neutralized phenol (100
ul), 0.5 M EDTA (100 ul) and 1% sodium dodecyl sulfate
(100 ul). The DNA was purified by gel filtration
chromatography through Sephadex G-100 followed by
25 precipitation with ethanol.

EXAMPLE V

Biotin and polybiotinylated poly-L-lysine were coupled to oligoribonucleotides using a carbodiimide coupling procedure described by Halloran and Parker, J. Immunol., 96 373 (1966). As an example, DNA (1 ug/ml), 1 ml) in tris buffer pH 8.2, sheared with 0.1 N sodium hydroxide was denatured by boiling for 10 minutes and quick cooling in an ice bath. Biotinyl-1,6-diaminohexane amide (2 mg, 6 umol) or polybiotinylated poly-L-lysine (2 mg) and 1-ethyl-3-diisopropylaminocarboimide HCl (10 mg, 64 umol) were added, and the pH readjusted to 8.2. After 24 hours at room temperature in the dark, the mixture was dialyzed against 10 mM tris buffered saline. DNA was precipitated ethanol.

EXAMPLE VI

Biotin, conjugated to cytochrome C, was prepared by the following procedure. To a solution of cytochrome C (10 mg) in 1 ml of 0.1 M sodium borate, pH 8.5 was added biotinyl-N-hydroxysuccinimide ester (10 mg, 29 umol) in 1 ml dimethyl formamide. After 4 hours at room temperature, the biotinylated protein was purified by gel filtration chromatography through a Sephadex G-50 column.

EXAMPLE VII

Formaldehyde coupling of cytochrome C-biotin and polybiotinylated poly-L-lysine to oligodeoxyribonucleotides were carried out using a method similar to that
05 described by Manning et al, Chromosoma, 53, 107 (1975). Oligodeoxyribonucleotide fragments obtained by sodium hydroxide shearing of purified DNA (100 ug/ml in 10 mM triethanolamine, pH 7.8 were denatured by boiling for 10 minutes followed by quick cooling in ice. Cytochrome C-
10 biotin 0.05 g/ml or polybiotinylated poly-L-lysine solution (0.05 ml) dissolved 3 mg/ml in 10 mM triethanolamine, pH 7.8 was added to 1 ml at the denatured oligodeoxyribonucleotide solution along with 0.1 ml of 6% formaldehyde in 10 mM triethanolamine, pH 7.8. After
15 stirring at 40° for 30 minutes the mixture was dialyzed against the same buffer. The oligodeoxyribonucleotide-biotin complex was finally purified by gel filtration chromatography on Sephadex G-100 followed by precipitation from ethanol.

EXAMPLE VIII

Double stranded polydeoxyadenylic acid:polybiotinylated deoxyuridylic acid was synthesized as follows. The double stranded oligonucleotide polydeoxyadenylic
05 acid:polythymidylic acid (20 ug) of length 300 basic pairs, dissolved in 200 ul exonuclease III buffer consisting of Tris-HCl pH 8.0 (70 mM); magnesium chloride (1.0 mM) and dithiothreitol (10 mM) was incubated with 100 units exonuclease III for 20 minutes
10 at 20°C. The partially digested oligonucleotide was immediately extracted with phenol, and the DNA was precipitated with 70% aqueous ethanol. The partially digested oligonucleotide was redissolved in 20 ul 5mM tris-HCl pH 7.6 and incubated at 20°C. for 2 hours in a
15 reaction containing 2'--deoxy- adenosine-5'-triphosphate (15 uM) thymidine-5'-triphosphate (the amount determines the degree of substitution) and biotinylated 5-(3-amino-1-propene) 2'-deoxyuridine-5'-triphosphate (5 uM), Klenow DNA polymerase I (200 units) dissolved in 0.1 mM
20 Potassium phosphate, pH 8.0 at a concentration of 0.2 units/ul. The biotinylated poly dA:poly dT, biotinyl dU was purified by gel filtration chromatography on Sephadex G-100. The DNA was ethanol precipitated and redissolved in 20 ul of solution containing sodium
25 acetate pH 4.6 (30 mM), sodium chloride (50 mM), zinc sulfate (1 mM) and glycerol (5%). S₁ nuclease (200 units) was added, and the reaction was incubated at 37° for 10 minutes. The reaction was stopped with 1 ml ammonium acetate (4 M) and 6 ml ethanol. The DNA was repurified by G-100 gel filtration chromatography and ethanol
30 precipitation.

EXAMPLE IX

Ligation of poly dA:poly dT, biotinyl dU to oligodeoxy-
ribonucleotides was accomplished as follows: DNA
fragments from alkali sheared purified DNA (as described
05 in Example VIII) were digested with S₁ nuclease and
repurified by phenol extraction and ethanol precipita-
tion. Blunt ended DNA fragments (1 ug) and poly dA:poly
dT, biotinyl dU (2 ug) were dissolved in 6 ul at a
buffer containing tris-HCl pH 7.4 (66 mM), magnesium
10 chloride (6.6 mM), adenosine triphosphate (24 mM) and
dithiothreitol (1.0mM), T₄ DNA ligase (50 units) was
added, and the volume brought to 20 ul with water. The
reaction was incubated 3 hours at 37°C. The DNA was
purified by gel filtration chromatography through
15 Sephadex G-100 and was ethanol precipitated.

EXAMPLE X

5-Hydroxymethyl-2'-deoxycytidylic acid was prepared by
20 enzymatic hydrolysis of non glycosylated phage T₄ DNA.
Purified phage DNA (2 mg), dissolved in 1 ml 50 mM tris
pH 7.4 and 10 mM magnesium chloride, was incubated 20
hours with deoxyribonuclease I at 37°. The pH was
adjusted to 9.0 and sodium chloride (20 mM) added.
25 Snake venom phosphodiesterase (0.05 g units in 0.5 ml
water) was added and incubation continued at 37° for 5
hours. An additional 0.05 units phosphodiesterase was
added and incubation continued 18 hours. Nucleotides
were separated by gel filtration chromatography through
30 Sephadex G-50. 5-hydroxymethyl-2'-deoxycytidylic acid
was purified by reverse phase high pressure liquid
chromatography.

EXAMPLE XI

5-(4-aminobutylaminomethyl)-2'-deoxyuridylic acid was obtained by enzymatic hydrolysis of DNA from phage ØW-
05 14. The phage was grown on Pseudomonas acidovorans 29 according to Kropinski and Warren, Gen. Virol. 6, 85 (1970), and the phage DNA purified according to Kropinski et al, Biochem. 12, 151 (1973). The DNA was enzymatically hydrolyzed with deoxyribonuclease I and
10 snake venom phosphodiesterase using the procedure described elsewhere (Example X). 5-(4-aminobutylamino-methyl)-2'-deoxyuridylic acid was purified by reverse phase high-pressure liquid chromatography.

EXAMPLE XII

Biotinylated-5-(4-aminobutylaminomethyl)-2'-deoxy-uridylic acid was prepared as follows: Biotinyl-n-hydroxysuccinimide ester (70 mg 0.2 m mol) dissolved in
20 1 ml dimethylformamide was added to 5-(4-aminobutyl-aminomethyl)-2'-deoxyuridylic acid in 20 ml 0.1 M sodium borate pH 8.5. After 4 hours the solution was concentrated to 0.5 ml by evaporation, and the biotinylated nucleotide was purified by reverse phase high pressure
25 liquid chromatography.

EXAMPLE XIII

5-formyl-2'-deoxyuridine prepared according to Mertes and Shipchandler, J. Heterocyclic Chem. 1, 751 (1970).

05 5-hydroxymethyluricil (1 mmol) dissolved in 20 ml dimethylsulfonate was heated at 100°C. with manganese dioxide (2.5 mmol) for 15 minutes. The solvent was evaporated at reduced pressure. The residue was taken up in hot ethanol and recrystallized from ethanol to yield 5-formyluracil, 5-formyluracil (0.10 g) was
10 silylated and dissolved in dry acetonitrile (2.5 ml). 2-deoxy-3,5-di-O-p-tolyl-D-ribofuranosyl chloride (Bhat,
Syn. Proc. in Nucleic Acid Chem., Vol. I, p. 521 (1968)
(0.22 g) and molecular sieves (0.2 g) were added, and
15 the mixture stirred at 25°C. for 40 hours under anhydrous conditions. The mixture was filtered and evaporated. The resulting oil was treated with anhydrous ethanol (2 ml) and chromatographed on silica gel to obtain the partially pure anomer which was re-crystallized from ethanol (M.P. 195-196°C.) The toluyl
20 groups were removed by reaction of the product in methanol benzene with sodium methoxide. The mixture was neutralized with Dowex 50 (H^+). 5-formyl-2'-deoxyuridine was recrystallized from ethanol M.P. 175-176°C.

EXAMPLE XIV

Biotin was coupled to 5-formyl-2'-deoxyuridine as follows:

To 5-formyl-2'-deoxyuridine (.320 g, 1.0 mmol) dissolved
in 300 ml 0.05 M sodium borate, was added biotinyl-1,6-

05 diaminohexane amide (0.74 g, 2 mmol). After stirring
one hour, sodium borohydride (0.2 g, 5 mmol) was added
and stirring continued for an additional 4 hours
followed by the addition of 8 ml 1M formic acid.

10 The biotinated compound was purified by reverse phase HPLC
eluting with methanol:0.5 M triethyl ammonium acetate, pH 4.0.

15

EXAMPLE XV

Biotin was coupled to 5-amino-2'-deoxyuridine as follows:

5-amino-2'-deoxyuridine (0.24 g, 1 mmol), biotin (0.25

20 g, 1 mmol) and dicyclohexylcarboimide (0.21 g, 1 mmol)
were dissolved in dry dimethyl formamide and stirred at
room temperature overnight. After filtration and
evaporation of the solvent, the residue was washed with
ether. The biotin-coupled product was purified by
25 reverse phase high pressure liquid chromatography using
a water methanol gradient.

EXAMPLE XVI

5-(oxy)acetic acid-2'-deoxyuridine was prepared according to a procedure of Deschamps and DeClerq, J. Med. Chem., 05 21, 228 (1978). 5-hydroxy-2-deoxyuridine (282 mg, 1.15 mmol) was dissolved in 1.16 ml, 1N potassium hydroxide (1.16 mmol) after which iodoacetic acid (603 mg, 3.4 mmol) in 1 ml water was added. After reaction at room temperature for 48 hours 1N HCl (1.06 ml) was added. 10 Concentration of this solution and addition of ethanol yielded a precipitate which was filtered, washed with cold ethanol and recrystallized from hot ethanol.

EXAMPLE XVII

15 Biotinyl-1,6-diaminohexane amide was coupled to 5-(oxy)acetic acid-2'-deoxyuridine as follows: Biotinyl-1,5-diaminohexane amide (0.74 g, 0.2 mmol), 5-(oxy)acetic acid-2'-deoxyuridine (0.60 g, 0.2 mmol) and dicyclohexylcarboimide (0.41 g, 0.2 mmol) were dissolved in 5 ml dry dimethylformamide and remained overnight at room temperature. The reaction was subsequently filtered and the solvent removed by evaporation. The residue was washed with .1N HCl and ether. The biotinated uridine derivative was purified by reverse phase high pressure liquid chromatography using a water-methanol gradient. 20 25

EXAMPLE XVIII

Phosphorylation of 5-substituted pyrimidine nucleosides was accomplished by the general procedure described below for biotinylated-5-(oxy)acetic acid-2'-deoxyuridine.

05 The nucleotide (0.16 g, 0.5 mmol) was dried by repeated evaporation from dry pyridine and redissolved in 10 ml dry pyridine. Monomethoxytrityl chloride (0.30 g, 0.8 mmol) was added and the mixture stirred at room

10 temperature in the dark for 18 hours. The solution was diluted with chloroform (200 ml) and extracted with 0.1 M sodium bicarbonate. The organic layer was dried and evaporated. The tritylated nucleoside was redissolved in dry pyridine (20 ml) and acetylated by reaction at room temperature with acetic anhydride (0.1 ml, 20 mmol). The mixture was cooled to 4°C. and methanol (40 ml) added. After stirring 10 hours at room temperature, the reaction was concentrated by evaporation. The compound was detritylated by dissolving in 1% benzene

15 sulfonic acid in chloroform (20 ml). After evaporation of solvent the nucleoside was purified by chromatography on silica gel eluting with 2% methanol:chloroform. The 3'-acetylated nucleoside was dried by repeated evaporation of dry pyridine. A mixture of phosphorous

20 oxychloride (100 ul, 1 mmol), (1-H), 1,2,4-triazoic (140 mg, 2.2 mmol) and triethylamine (260 ul, 2.0 mmol) was stirred in 5 ml anhydrous dioxane at 10°-15°C. for 30 minutes and at room temperature for 1 hour. This was added to the 3'-acetylated nucleoside, and the mixture

25 stirred at room temperature for 1 hour after which it was cooled to 0°C. Water (5 ml) was added and the reaction stirred at room temperature for 18 hours. Barium chloride (100 mg, 5 mmol) was added and the barium salt of the nucleotide collected by filtration.

30 The salt was washed with water and ether. The barium

35

salt was converted to the sodium salt by stirring with Dowex 50 (Na^+ form) in 10 ml water for 4 hours at room temperature. 2 N sodium hydroxide (2N, 10 ml) was added and the reaction stirred for 15 minutes at room temperature after which it was neutralized by addition of excess Dowex 50 (H^+) form. The deacetylated nucleotide was concentrated by evaporation and purified by reverse phase high-pressure chromatography.

10 EXAMPLE XIX

5-substituted pyrimidine triphosphates were chemically prepared from their respective 5' monophosphates using a procedure of Michelson, Biochem Biophys Acta, 91, 1, (1964). The example of 5-hydroxymethyl-2'-deoxycytidine-5'-triphosphate will be given. The others were similarly prepared. 5-hydroxymethyl-2' deoxycytidyllic acid (free acid) (0.63 g, 0.2 mmol) was converted to its tri-n-octylammonium salt by suspending in methanol and addition of tri-n-octylammonium hydroxide (0.74 g, 0.2 mmol). The suspension was refluxed until a clear solution was obtained and the solvent removed under vacuum. The salt was dried by dissolution in and subsequent evaporation from dry pyridine several times. To the salt, dissolved in dry dimethylformamide (0.1 ml) and dioxane (1 ml) was added diphenylphosphochloride (0.1 ml) and tri-n-butylamine (0.2 ml). After 25 hours at room temperature, solvent was removed and ether was added to precipitate the nucleoside-5'-diphenylpyrophosphate. This was dissolved in dioxane (0.5 ml) and a solution of di(tri-n-butylammonium) pyrophosphate (0.5 mmol) in 1 ml pyridine was added. After 45 minutes at room temperature, the mixture was concentrated under

vacuum to a small volume. The crude product was precipitated with ether. This was dissolved in 0.1 M phosphate buffer pH 8.0. The trisphosphate was purified by chromatography on DEAE cellulose eluting with a
05 gradient of 0.1 to 0.6 M triethylammonium bicarbonate ph 7.5.

EXAMPLE XX

10 DNA was labeled with 5-substituted pyrimidine triphosphates by nick translating DNA in the presence of the appropriate triphosphate. An example follows for labeling purified DNA with biotinylated 5-formyl-2'-deoxyuridine. DNA (20 ug/ml) was incubated at 14°C. in the presence of magnesium chloride (5 mM) 2'-deoxy-
15 cytidine-5'-triphosphate (15 mM), 2'-deoxyadenosine-5'-triphosphate (15 uM), 2'-deoxyguanosine-5'-triphosphate (15 uM), biotinylated-5-formyl-2'-deoxyuridine-5'-triphosphate (20 uM), activated pancreatic deoxyribonuclease I (13 mg/ml), E. coli deoxyribonuclease acid,
20 polymerase I (40 units/ml) and tris HCL, pH 7.4 (50 mM). After 2 hours the reaction was stopped by addition of 0.3 M EDTA (.05 ml) followed by heating at 65° for 5 minutes. Labeled oligonucleotide was purified by gel filtration chromatography through Sephadex G-100 and
25 precipitation from cold ethanol.

EXAMPLE XXIPRECIPITATION OF GLUCOSYLATED DNA
BY CONCANAVALIN A

05

Reaction mixtures (1.0 ml) were prepared in 1.5 ml eppendorf tubes as follows:

	Sodium potassium phosphate, pH 6.5	10mM
	NaCl	150mM
10	MgSO ₄	5mM
	CaCl ₂	1mM
	DNA (T4 of calf thymus)	50ug
	Concanavalin A (10 mg/ml)	50-500 ug

15

Reactions were started by the addition of concanavalin A (Con A). The solutions were mixed and left at room temperature for 60 minutes. The tubes were centrifuged at 1200 g for 15-20 minutes. The supernatants were diluted and the A₂₆₀ was measured.

20

Since Con A absorbs at 260 nanometers, control solutions lacking DNA but containing Con A were prepared. The Con A absorbance was subtracted from the absorbence obtained in the complete reaction mixtures.

25

The results of this reaction are presented in accompanying Figure 1.

EXAMPLE XXIIBINDING OF GLUCOSYLATED DNA
TO CONCANAVALIN A

05 Phage T4 DNA and phage DNA were labeled by incorporation of H^3 -deoxyadenosine triphosphate into the DNA by nick translation according to the Rigby et al procedure. T4 DNA was nick translated to a specific activity of 5×10^5 cpm/microgram and an average double- stranded size of 5 kilobases. Lambda DNA was nick translation to a specific activity of 3×10^5 cpm/microgram and an average double stranded size of 6.0 kilobases as determined by agarose gel electrophoresis. Unincorporated nucleotides were removed from the reaction mixtures by Bio-Gel P-60 chromatography.

15 Con A sepharose was prepared as described by the manufacturer (Pharmacia). One ml of settled gel contained 18 mg of bound Con A. One ml columns were prepared in sterile pasteur pipettes and were 20 equilibrated with PBS (0.15 M NaCl; .01 M sodium potassium phosphate, pH 6.5).

25 H^3 -DNA samples were prepared in 0.5 ml of buffer (as described in Example XXI but without Con A). T4 DNA solutions contained 176,000 cpm/0.5 ml, and DNA solutions contained 108,000 cp./0.5 ml. A 0.5 ml sample was applied to the column.

30 A 10.5 ml volume of buffer was passed through the column, and the eluate fractions (0.33 ml) were collected and counted in a Beckman LSC-100 scintillation counter in a 3.5 ml reafluor cocktail (Beckman). The results

(Figures 2A and 2B) show that non-glucosylated DNA was not bound whereas glucosylated T4 DNA was bound to the column. The bound T4 DNA was removed by washing the column with a higher pH buffer (Tris-HCl, pH 7.2 - 8.2).

05

Furthermore, consistent with the interaction of glucose and Con A, mannose, when included in the buffer in which the DNA is applied to the column, prevents binding of glucosylated DNA to Con A sepharose. Also, mannose-containing buffer (PBS-containing 0.056 M mannose) removes bound T4 DNA from Con A sepharose (Figures 3A and 3B).

10

15 Further illustrative of the practices of this invention directed to nonradioactive methods or techniques of assaying for specific nucleic acids, the following example deals with the use of the sugar-lectin system. This example deals with the use of DNA which is not glycosylated in nature but rather has had a maltotriose group added thereto by way of nick translation described herein. The maltotriose modified dUTP and DNA modified therewith bind specifically to a column of concanavalin A covalently bound to sepharose. By this technique and in accordance with the practices of this invention, there 20 is provided a means for specifically labeling any nucleic acid with sugars. As previously indicated herein, nick translation is only one of a number of techniques and approaches possible for the production of 25 the modified nucleic acids in accordance with this invention.

25

30

EXAMPLE XXIII

Lambda DNA was nick translated as described herein with maltotriose coupled to 5-(3-amino-1-propenyl)-2'-deoxy-
05 uridine-5' triphosphate and ^{3}H -2'-deoxyadenosine-5'-triphosphate. Under these conditions DNA was substituted to 40 per cent of its thymidine residues with the maltotriose nucleotide and had a specific activity of 8×10^5 counts per minute (cpm) per microgram
10 of DNA. A control sample of DNA substituted only with ^{3}H -dATP had a specific activity of 6×10^5 cpm per microgram DNA. The nick translated DNA samples were purified free of reaction mixture components by Biogel P-60 chromatography as described herein.

15 The purified samples were applied to Con A-sepharose columns as described in Figures 2A and 2B. The maltotriose-labeled DNA was retained on the column when washed with PBS but was removed by subsequent elution
20 with 10mM Tris-HCl, pH 8.2 (Figure 4A). The unsubsti-
tuted tritiated DNA did not bind to the column at pH 7.4 (Figure 4B).

EXAMPLE XXIV

Potentially immunogenic heptenes may be introduced at the 5 position of uridine by a variety of methods in the literature. 5-(perfluorobutyl)-2'-deoxyuridine was synthesized using a method of Cech et al, Nucl. Acids Res. 2, 2183 (1979). Copper-bronze was prepared by reacting copper sulfite (5 g, 20 mmol) with zinc powder (2 g) in 20 ml water. The mixture was decanted, and the residue washed with water and then 5% hydrochloric acid and water. Just before use, the solid (2 g) was activated with 2% iodine in acetone (20 ml). After filtration the residue was washed with acetone:concentrated hydrochloric acid and then pure acetone.

Activated copper-bronze (130 mg, 2 mmol) and 1-iodo-1',2,2',3,3',4,4'heptafluorobutane (1.3 mg, 4 mmol) were stirred in 3 ml dimethylsulfoxide at 110°C. for 1 hour. After cooling and filtration, 2'-deoxyuridine (245 mg, 1mmol) was added, and the mixture heated at 110°C. for 1 hour. Water (5 ml) was added, and the mixture extracted with ether. The ether extracts were dried and evaporated under reduced pressure. The residue was chromatographed on a silica gel column eluting with ethylacetate.

25

EXAMPLE XXV

Tubericydin was substituted at the 5 position by derivitizing the 5-cyano compound, toyocamycin. An example is the synthesis of 4-amino-5 (tetrazol)-5-yl)-7- (B-D-ribofuranosyl) pyrrolo[2,3-d]pyrimidine using a procedure of Schram and Townsend, J. Carbohydrate, Nucleosides:Nucleotides 1, 38 (1974). Toyocamycin (1.0 g) dissolved in water (100 ml) and glacial acetic acid (13 ml) was heated to reflux. Sodium azide (7.5 g) was added in 1.25 g portions over 10 hours. The solution was cooled to 5°C. and the precipitated product collected, M.P. 276-277°C.

EXAMPLE XXVI

5-Cyano-2'-deoxyuridine was prepared according to Bleckley et al, Nucl. Acids Res. 2, 683 (1975). 5-Iodo-
05 2'-deoxyuridine (1.0 g, 2.82 mmol) was dissolved in refluxing hexamethyldisilizane (HMDS) (10 ml). Excess HMDS was removed at reduced pressure, and the resulting oil was dissolved in dry pyridine (50 ml). Cuprous cyanide (350 mg, 3.8 mmol) was added, and the solution
10 heated at 160°C. for 20 hours. Pyridine was removed at reduced pressure, and the residue extracted into toluene which was subsequently evaporated. The residue was heated in 50% aqueous ethanol at 100° for 2 hours. The product was purified by reverse-phase high pressure
15 liquid chromatography and recrystallized from ethanol, M.P. 161°C.

EXAMPLE XXVII

20 4-amino-5-amino methylene-7-(β -D-2-deoxyfuranosyl) pyrrolo[2,3-d]pyrimidine dihydrochloride was obtained as follows. 4-amino-5-cyano-7-(β -D-2-deoxyfuranosyl) pyrrolo[2,3-d]pyrimidine (Toyocamycin) (0.2 g) was dissolved in hydrochloric acid (10 ml). 10% palladium
25 on charcoal (0.1 g) was added as the mixture hydrogenated at 40 psi for 5 hours at room temperature. After filtration the water was evaporated at reduced pressure. The residue was triturated with ethanol, and the product recrystallized from 50% ethanol.

EXAMPLE XXVIII

5-amino-2'-deoxyuridine was prepared from 5-bromo-2'-deoxyuridine according to the procedure of Roberts and Visser, J. Am. Chem. Soc. 14:665-669 (1952). 5-bromo-
05 2'-deoxyuridine (2g, 6.2 mmol) dissolved in liquid ammonia (20 ml) was sealed in a glass tube and heated at 50° for 5 days. The tube was opened, and the ammonia was evaporated. 5-amino-2'-deoxyuridine was recrystallized from 5 ml water and 75 ml hot isopropyl alcohol.

10

EXAMPLE XXIX

5-(methylamino)-2'-deoxyuridine (0.2 g) was prepared as follows. 5-cyano-2'-deoxyuridine (0.2 g, 0.05 mol) was
15 dissolved in 1 N hydrochloric acid (10 ml). 10% palladium on charcoal (0.1 g) was added, and the mixture hydrogenated at 40 p.s.i. for 10 hours at room temperature. The mixture was filtered and the water evaporated at reduced pressure. The residue was triturated with
20 ether, and the product was recrystallized from 80% ethanol.

EXAMPLE XXX

25 Maltose triose was oxidized to the corresponding carboxylic acid by the following method. Maltose triose (0.5 g, 0.94 mmol) was dissolved in water (5 ml). Lead carbonate (0.42 g, 1.1 mmol) and bromine (0.17 ml), 3.3 mmol) were added, and the mixture was allowed to react
30 at room temperature for six days after which no reducing sugar remained. The mixture was filtered, and silver carbonate (0.2 g) added. After refiltering, the filtrate was deionized by elution through Dowex 50 (H^+ form). Evaporation of water and drying in the presence
35 of phosphorus pentoxide yielded the desired product.

0097373

EXAMPLE XXXI

Maltose triose was coupled to 5-(3-amino-1-propenyl)-
2'-deoxyuridine-5'-triphosphate by the following
05 procedure. Oxidized maltose triose (190 mg, 0.18 mmol)
was dissolved in dimethylformamide (0.8 ml) and cooled
to 4°C. Isobutyl chloroformate (25 mg, 0.18 mmol) and
tri-n-butylamine (43 ul, 0.38 mmol) were added, and the
solution allowed to react at 4°C. for 15 minutes. 5-(3-
10 amino-1-propenyl)-2'-deoxyuridine-5'-triphosphate (9.0
μmol), dissolved in dimethyl formamide (1.2 ml) and 0.1
M sodium borate and cooled to 4°C., was added to the
above solution. The mixture was incubated at 4°C. for 1
hour and at room temperature for 18 hours. It was then
15 loaded on a DEAE-cellulose column and eluted with a
gradient of 0.1 to 0.6 M triethylammonium bicarbonate,
pH 7.5. The product was finally purified by reverse
phase high pressure liquid chromatography.

20 Following are Examples XXXII and XXXIII. Example XXXII
is a method of tagging allylamine modified dUTP with a
fluorescein substituent. This is an example of creation
of a self detecting nucleic acid probe. Example XXXIII
25 is a method of labeling preformed double helical nucleic
acids at the N² position of guanine and the N⁶ position
of adenine. Example XXXVII has the detector molecule
linked to the probe. Chromosoma 84: 1-18 (1981) and
Exp. Cell Res. 128:485-490, disclose end labeling of RNA
with rhodamine. However, the procedure of this
30 invention is less disruptive and labels internal nucleo-
tides.

EXAMPLE XXXII

Fluorescein was coupled to 5-(3-amino-1-propyl)-2'-
deoxyuridine-5'-triphosphate (AA-dUTP) as follows. AA-
05 dUTP (10 umol), dissolved in 2 ml sodium borate buffer
(0.1 m); pH 9.0, was added to fluorescein isothiocyanate
(10 mg, 25 umol) dissolved in 1 ml dimethylformamide.
After four hours at room temperature the mixture was
loaded onto a DEAE-cellulose column equilibrated in
10 triethylammonium bicarbonate buffer, pH 7.5. The
fluorescein coupled AA-dUTP was purified by elution with
a gradient of from 0.1 to 0.6 m triethylammonium
bicarbonate, pH 7.5.

EXAMPLE XXXIII

DNA may be modified by reaction with chemical alkylating
agents. Lambda DNA was alkylated in N² position of
guanine and N⁶ position of adenine by reacting DNA with
aromatic hydrocarbon 7-bromomethylbenz[a]anthracene. 7-
20 bromomethylbenz[a]anthracene was obtained as follows. 7-
methyl[a]anthracene in carbon disulfide solution was
cooled in a freezing mixture and treated dropwise with a
molar equivalent of bromine. After 30 minutes, the
25 product in suspension was collected, and was washed with
dry ether and recrystallized from benzene. The yield
was 66% with melting point 190.5-191.5°C.

DNA, purified from phage Lambda, (1.6 mg) was solubil-
30 ized in 5.0 ml of 20 mM potassium phosphate pH 6.5. To
4.0 ml of DNA solution was added 500 micrograms 7-
bromomethylbenz[a]anthracene in dry acetone. After 30
minutes at 20°, the DNA was precipitated with two
volumes of cold ethanol. The precipitate was washed
35 successively with ethanol, acetone and ether to remove

any unbound 7-bromomethylbenz[a]anthracene. Enzymatic hydrolysis of the DNA to nucleosides and subsequent chromatography of the products on Sephadex LH-20 columns, indicated that 18% of the adenine and 48% of
05 the guanine in DNA were modified in N⁶ and N² positions, respectively.

The modified DNA was made single stranded either by (1) heating to 100° for 5 minutes and rapid cooling or (2)
10 incubating with equal volume of 0.1 M NaOH for 10 minutes and then dialyzing the solution for four hours against 1 ml tris-HCl pH 8.0 containing 0.5 ml EDTA to keep the DNA in single-stranded form.

15 EXAMPLE XXXIV

A DNA probe was ligated to a synthetic DNA composed of repeated sequences of E. coli lac operator DNA. After hybridization to detect antiprobe sequences, the
20 hybridized DNA was detected by reaction with biotinylated lac repressor which was, in turn, detected by an enzyme linked immuno sorbent assay using goat antibiotin IGG to react with the biotin and a second antibody coupled to horse radish peroxidase. The lac polyoperator DNA has been described by Caruthers (Second Annual
25 Congress for Recombinant DNA Research, Los Angeles, 1982), and it was ligated, in a blunt end ligation, using T4 ligase, to an adenovirus DNA probe. In situ hybridization of the polyoperator-labeled probe DNA was carried out as described by Gerhard et al (Proc. Natl. Acad. Sci. USA, 78, 3755 (1981)). Biotinylated lac
30 repressor was prepared as described by Manning et al (Chromosoma, 53, 107-117 (1975)) and was applied to adenovirus infected cells, fixed to a glass slide, in Binding buffer composed of (0.01 M KCl, 0.01 M tris (pH
35

7.6), 0.01 M MgSO₄, 10⁻⁴ M EDTA, 10⁻⁴ M DTT, 5% DMSO (dimethyl sulfoxide) and 50 ug/ml bovine serum albumin by J. Miller, Experiments in Molecular Genetics, Cold Spring Harbor Laboratory (1972). The slides were washed in binding buffer to remove unbound biotinylated lac
05 repressor and then assayed for biotin using the horse radish peroxidase-linked double antibody procedure. This procedure could be adapted to create an affinity column where the probe could be bound to immobilized repressor protein and then removed by elution with a
10 specific inducer, for example, isopropylthigalactoside or thiomethylgalactoside. The affinity of the repressor-operator complex is quite high 10⁻¹¹ M. When a specific inducer binds to the repressor the operator-repressor complex collapses.

15

EXAMPLE XXXV

20 5-Bromo-2'-deoxyuridine-5'-phosphate was prepared as follows: 2'-Deoxyuridine-5'-phosphate (6.2 g) was suspended in a mixture of 60 ml pyridine and 30 ml acetic acid. Bromine (0.84 ml) was added with stirring in an ice water bath and stirring continued for 20 hours at room temperature. The solution was concentrated by vacuum. After redissolution in a minimum of water a
25 crude product was precipitated by addition of ethanol. The crude product was chromatographed on Dowex 50 (H⁺) and eluted with water. The free acid product was precipitated from the concentrated eluent by addition of ethanol.

30

EXAMPLE XXXVI

Calf intestine alkaline phosphate was biotinylated as follows: The enzyme (1 mg, 7.7 mmol) was chromatographed on a G-50 column eluting with 0.1 M Hepes buffer pH 8.0 containing 0.1 M sodium chloride. The pooled fractions were reacted with N-biotinyl-6-amino-caproic acid-N-hydroxysuccinimide ester (0.675 mg, 0.77 umol) dissolved in 10 ml diethylformamide at room temperature for 1 hour. Sodium periodate (0.1 M 125 ul) was added and stirring continued for 2 hours. The mixture was dialyzed at 4° overnight in 0.1 M Hepes buffer pH 8.0 with 0.1 M NaCl after which the pH was adjusted to 7.4. Biotin hydrazide (0.1 M, 0.5 ml) dissolved in 0.1 M Hepes buffer pH 7.4 and 0.1 M NaCl was added and the reaction stirred for 30 minutes at room temperature. The pH was adjusted to 8.0 with 0.2 M sodium carbonate and 0.5 ml of freshly prepared 0.1 M sodium borohydride in water was added, the solution was dialyzed against 0.1 M tris buffer pH 8.0 with 0.1 M NaCl.

20

EXAMPLE XXXVII

6-Cyano-2'-deoxyuridine-5'-phosphate was prepared similarly to a procedure of Veder et al, J. Carbohydr. Nucleosides, Nucleotides, 5, 261 (1978). 5-bromo-2'-deoxyuridine-5'-phosphoric acid (2.0 g, 15 mmol) dried by successive evaporation from pyridine was dissolved in 50 ml dimethylsulfide. Sodium cyanide (490 mg, 10 mmol) was added and the solution stirred at room temperature for 2 days. The solution was diluted with 400 ml water and the pH adjusted to 7.5. It was applied to a DEAE-cellulose column (HCO^-_3 form) washed with 2000 ml 0.02 M triethylammonium bicarbonate to yield the desired product.

EXAMPLE XXXVIII

6-(Methylamino)-2'-deoxyuridine-5'-phosphoric acid was prepared as follows: 6-Cyano-2'-deoxyuridine-5'-phosphoric acid (0.2 g, 60 mmol) was dissolved in 0.1 M hydrochloric acid. After addition of 10% palladium on charcoal (0.1 g), the mixture was hydrogenated at 40 psi for 20 hours at room temperature. The mixture was filtered, neutralized with lithium hydroxide and lyophilized. The product residue was extracted with ethanol.

EXAMPLE XXXIX

Horse radish peroxidase (20 mg) dissolved in 5 ml distilled water was added to 1.0 ml freshly prepared 0.1 M sodium periodate solution. After stirring at room temperature for 20 minutes it was dialyzed overnight at 4°C. against 1 mM sodium acetate pH 4.4. Biotin hydrazide (2.6 mg, 5x10⁻² mmol) dissolved 2.0, 0.1 M Hepes buffer pH 7.4 with 0.1 M sodium chloride was brought to pH 8.0 with 0.2 M sodium carbonate and 0.5 ml of a freshly prepared 0.1 M sodium borohydride solution in water was added. After 2 hours at 4°C. the protein was purified on a Sephadex G-50 column eluting with 0.1 M Hepes and 0.1 M NaCl.

EXAMPLE XL

Carrot acid phosphatase has been mentioned by Brunngraber and Chargaff, J. Biol Chem., (1967) 242, 4834-4840 as a byproduct of the purification of phosphotransferase and has been purified to a specific activity of 460 uM/mg/min

at 37°C. with paranitrophenylphosphate as the substrate. The purification involved the steps of (a) absorption of non-specific proteins by DEAE cellulose; (b) acid purification of the enzyme; (c) acetone fractionation; 05 (d) concanavalin A affinity chromatography; (e) hydroxyapatite chromatography and (f) Sephadex G-100 fractionation. The specific activity of the enzyme subjected to the Sephadex G-100 fractionation due to loss of activity in the preceding affinity chromatography step (d) was 10 170 uM/mg/m. By changing elution conditions at step (d), these losses can be avoided with the result that the specific activity of the enzyme before the Sephadex G-100 fractionation can be improved to 340 uM/mg/m. The Sephadex G-100 fractionation step should yield an enzyme 15 having a specific activity of 800 uM/mg/m or higher.

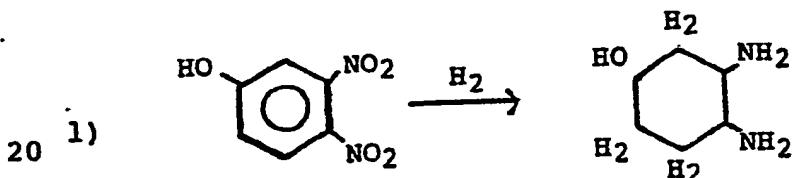
Carrot acid phosphatase was biotinylated using a procedure of Wilchek et al Biochemistry 6, 247 (1967). To the enzyme (20 mg) dissolved 0.1 M NaCl, pH 5, was added biotin hydrazide (2.0 mg, 7×10^{-3} mmol) and 1-ethyl-3-(3-dimethylaminopropyl) carbodimide hydrochloride 20 (1 mg, 7×10^{-3} mmol) dissolved in 0.1 M NaCl, pH 5. After 2 hours at 4°C. the enzyme was chromatographed on Sephadex G-50 eluting with 0.1 M sodium acetate, pH 5.0.

Of special importance and significance in the practices of this invention is the utilization of self-signaling or self-indicating or self-detecting nucleic acids, particularly such nucleic acids which are capable of

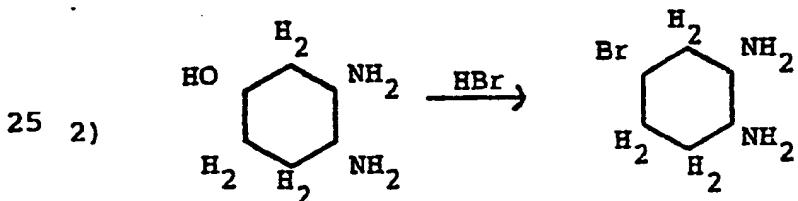
05 being incorporated in double-stranded DNA and the like. Such self-signaling or self-detecting nucleic acids can be created by covalently attaching to an allylamine substituent making up a modified nucleotide in accordance with this invention a molecule which will chelate

10 specific ions, e.g. heavy metals, rare earths, etc. In general, the chelated ion can be detected either (a) by radioactive emission or (b) by using the ion to catalyze a chromogenic or fluorogenic reaction.

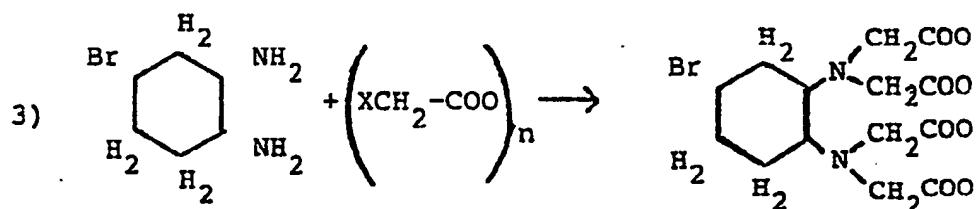
15 By way of example, a solution of 3,4-dinitro phenol is reduced to 3,4-diamino cyclohexane



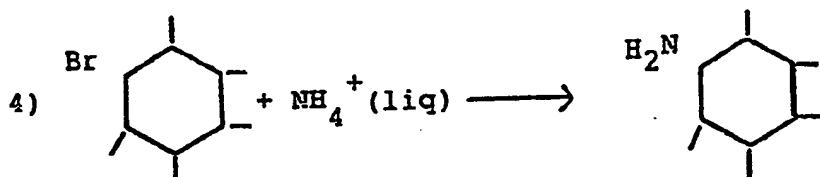
This material is then brominated



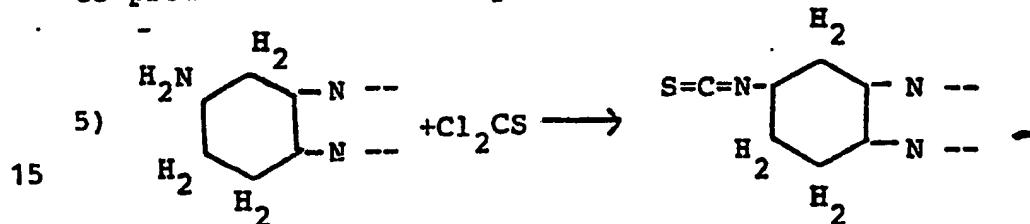
to form 3,4-diamino bromo cyclohexane (dABCH). This compound is reacted with halide (Cl, Br, I) substituted
30 carboxymethyl compound to produce a tetra carboxymethyl derivative or dABCH (TCM-dABCH):



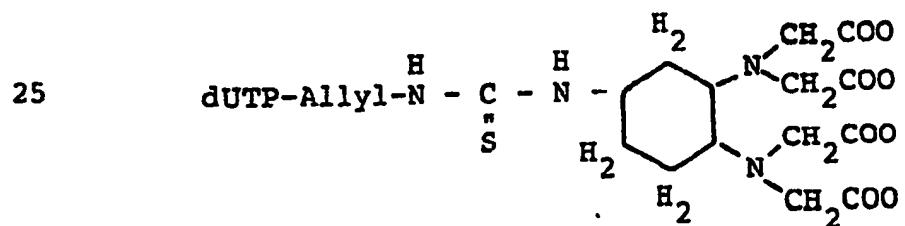
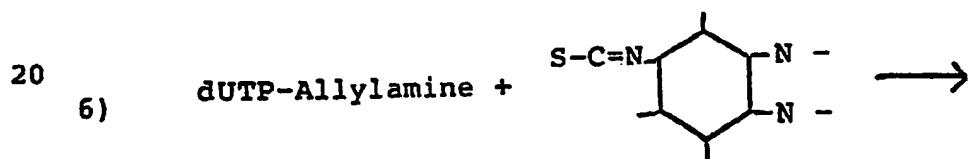
05 The bromine is substituted by an amino group using soluble ammonia:



10 Then this compound is reacted with chloro thiophosgene to produce the isothiocyanate derivative of (TCM-dANCH).



Finally, this compound is reacted with dUTP-allylamine derivative to produce modified dUTP.



30 Cobalt or other heavy metal ions or other rare earth ions can be chelated to the compound after step 3 above. Or the nucleic acid can be substituted with this adduct and then the ion added. (Example, cobalt is added at pH 6 where the binding constant is 10⁻¹⁹M).

Cobalt can be assayed by radioactivity. It can also be detected by its ability to oxidize methylene blue to the leuco form in the presence of molecular oxygen. It can be used to oxidize soluble sulfhydro groups to disulfide bonds again in the presence of molecular oxygen.

This type of self-signaling molecule can be used to monitor any nucleic acid hybridization reaction. It is particularly important for detecting nucleic acids in gels (for example, sequencing gels).

With respect to its use in radioactivity, it can be used to tailor the isotope needed, i.e. if a weak or strong β or γ emitter is needed, that isotope can be chelated. Examples of isotopes that can be used are listed immediately hereinafter.

Antimony-124	Iodine-125	Scandium-44
Antimony-125	Iodine-131	Scandium-46
Arsenic-74	Iodine-132	Selenium-75
	Iridium-192	Silver-110m
Barium-133	Iron-55	Silver-111
Barium-140	Iron-59	Sodium-22
Beryllium-7	Krypton-85	Strontium-85
Bismuth-208	Lead-210	Strontium-89
Bismuth-207	Lutecium-177	Strontium-90
Cadmium-109	Manganese-54	Sulphur-35
Cadmium-115m	Mercury-197	
Calcium-45	Mercury-203	Tantalum-182
Carbon-14	Molybdenum-99	Technetium-99
Cerium-139	Neodymium-147	Tellurium-125m
Cerium-141	Neptunium-237	Tellurium-132
Cerium-144	Nickel-63	Terbium-160
Cesium-134	Niobium-95	Thallium-204
Cesium-137	Osmium-185+191	Thorium-228
Chlorine-36	Palladium-103	Thorium-232
Chromium-51	Platinum-195m	Thulium-170
Cobalt-56	Praseodymium-143	Tin-113
Cobalt-57	Promethium-147	Titanium-44
Cobalt-58	Protactinium-233	Tritium
Cobalt-60	Radium-226	Tungsten-185
	Rhenium-186	
Erbium-169	Rubidium-86	Vanadium-48
Europium-152	Ruthenium-103	Vanadium-49
Gadolinium-153	Ruthenium-106	Ytterbium-169
Gold-195		Yttrium-88
Gold-199		Yttrium-90
Hafnium-175		Yttrium-91
Hafnium-175+181		Zinc-65
Hafnium-181		Zirconium-95
Hydrogen-3 see Tritium		

Streptavidin, a protein produced by a Streptomyces avidinii is a large molecular weight component of a synergistic pair of compounds which are both present in the culture filtrates of this microorganism. Each of
05 the pair is inactive but in combination are active against gram-negative microorganisms. It has been found that the small component of this antibiotic prevents de novo synthesis of the vitamin biotin and thus, at least in synthetic media, show antimicrobial activity. In
10 complex medium, however, the large component has to be included to exert the same effect on bacteria. This has been shown to be due to the presence of external biotin in the complex medium. The large molecular component has been found to bind external biotin and thus demonstrating the same kind of action as avidin from eggs and
15 oviduct tissues of laying birds.

Streptavidin has been purified and shown to be a 60,000 dalton polypeptide. Like avidin, streptavidin contains
20 four subunits and binds tightly four molecules of biotin. Unlike avidin, however, it is non-glycosylated and it has PI of 5.0 as compared to avidin with PI = 10.5. Due to the difference in pI streptavidin does not have a tendency to non-specifically interact with DNA.

25 PREPARATION OF STREPTAVIDIN

A semi-synthetic medium containing salt, 1% glucose, 0.1% asparagine, 0.05% yeast extract and trace elements
30 was prepared. The cultures were grown at 26°C. for three days. Mycellium was removed by centrifugation and protein in the supernatant were absorbed to DEAE-cellulose in a batchwise process after pH was adjusted with 1M HCl to 7.2. DEAE-cellulose was filtered off and
35 washed with 20 mM Tris-HCl (pH 7.2) until no absorbancy

at 280 nm was recorded. Streptavidin was eluted with 20 mM Tris-HCl (pH 7.2) containing 0.5 M NaCl. Ammonium sulfate precipitation was used to further concentrate the streptavidin (50% w/v at 4°C.).

05

The precipitate was dissolved in water and dialyzed against 1.0 M NaCl, 50 mM Na₂CO₃. In the next step affinity column chromatography on iminobiotin sepharose was used. Eluted streptavidin from iminobiotin sepharose column was shown to be chromatographically pure by non-denaturing agarose-gel electrophoresis.

15

The final purification of streptavidin is accomplished by affinity purification through an iminobiotin-sepharose column. Iminobiotin is an analog of biotin in which the carbonyl of the urea moiety is substituted with an imine function. Iminobiotin will bind avidin and streptavidin at basic pH but the complex is dissociable at acidic pH.

20

Iminobiotin is prepared from biotin in several steps. Biotin is hydrolyzed by barium hydroxide to cis-3,4-diamino-2-tetrahydrothiophene-valeric acid which is reacted with cyanogen bromide to iminobiotin. The iminobiotin is coupled to amino sepharose via the N-hydroxysuccinimide ester of its hydrobromide salt.

25

The crude protein mixture from DEAE eluted Streptomyces avidinii incubation media is dissolved in 50 mM sodium carbonate and 1.0 M sodium chloride (pH 11) and applied to an iminobiotin column pre-equilibrated with this solution. The column is eluted at pH 11. Streptavidin is subsequently eluted with 50 mM ammonium acetate, pH 4.0 containing 0.5 M sodium chloride. The eluent is dialyzed three times against 1 mM Tris pH 7.4 and lyophilized to dryness.

35

In the practices of this invention avidin is useful as a detecting mechanism for labeled DNA, such as biotin-labeled DNA. However, avidin itself, such at about neutral pH, complexes with DNA with the result

05 that any signal derivable from a complex between biotin-labeled DNA and avidin might be lost or be non-detectable in the background due to the complex formation between avidin and unlabeled DNA. This disadvantage of the use of avidin in the practices of this invention is

10 not possessed by streptavidin which does not form a complex with DNA at about neutral pH but is capable of forming a complex with the biotin portion of biotin-labeled DNA.

15 In another aspect directed to the broad utility of avidin and streptavidin for detecting labeled compounds other than DNA, avidin and streptavidin are particularly effective as detecting mechanisms for labeled proteins, polysaccharides and lipids. By way of example, one can

20 fix to a solid matrix a specific antigen and bind to this antigen an antibody directed against this antigen which itself has been biotinylated. Then one can assay for the presence of this biotinylated antibody by reacting it with avidin or streptavidin complexed with an enzyme, such as calf intestine alkaline phosphatase, or to which

25 fluorescing molecule, as for example fluorescein has been linked.

The use of the antigen-antibody system for detecting either antigen or antibody is well known. A comparable

30 system is a system based on a glycosylated substrate or molecule and matching or appropriate lectin. In this system the lectin would carry a label, such as fluorescein or appropriate enzyme. In this glycosyl-lectin system

35 the labeled lectin forms a complex with the glycosyl

moiety, comparable to the antigen-antibody complex, and this complex comprising the glycosylated molecule and appropriate labeled lectin having the necessary glycosyl or sugar moiety specificity would then present
05 itself eliciting the expected response from the label portion of the labeled lectin making up the glycosyl-lectin complex.

Another aspect of the practices of this invention which
10 is particularly advantageous is to carry out the detection or hybridization in the liquid phase between the DNA sought to be detected and the DNA detecting probe. In this liquid phase system both the DNA molecule to be detected and the appropriate DNA detecting probe are not
15 attached to any insoluble substrate or any insoluble chemical moiety. The advantages of the liquid phase detection system reside in the speed of hybridization or hybrid formation between the DNA to be detected and the appropriate DNA probe therefor. For example, in
20 a solid-liquid system the time required to effect recognition and hybridization formation is about ten times greater than if it were carried out in a completely liquid system, i.e. both DNA to be detected and the detecting DNA are not attached to an insoluble moiety.

25 The probes prepared in accordance with the practices of this invention are adaptable for use in the detection of viruses and bacteria in fluids as indicated hereinabove. Where the fluids to be examined do not contain
30 large amounts of protein, the viruses therein can be concentrated by absorption on hydroxyapatite and eluted in a small amount of phosphate buffer. When the fluid to be examined contains large amounts of protein, the viruses can be concentrated by high speed centrifugation.

If antibody were available, absorption on an affinity column and elution with acid would be preferable because it would be possible to process many probes in accordance with the practices of this invention at the same time.

05 The bacteria to be examined is usually readily concentrated by centrifugation.

In accordance with the practices of this invention, the identification or characterization of the isolated particles, viruses and bacteria, would be hybridization of the characterizing or identifying DNA thereof with a specific single stranded DNA probe prepared in accordance with the practices of this invention. After hybridization, excess non-hybridized probe DNA would be digested with 15 S₁ nuclease and exonuclease I from E. coli at high salt content to suppress the nicking activity of the S₁ nuclease, see Vogt, Methods in Enzymology, Vol. 65, pages 248-255 (1980). This nuclease treatment would produce mononucleotides from the excess, non-hybridized 20 single-stranded DNA probe but would leave the double-stranded, hybridized DNA intact. This would then be absorbed at high salt content on Dowex anion exchanger (the nucleotides and the small amount of oligonucleotides will not bind to the resin in high salt concentration). The resulting hybridized DNA would then be 25 identified or characterized by various procedures applicable to the practices of this invention.

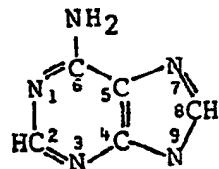
The special nucleotides of this invention include a phosphoric acid P moiety, a sugar or monosaccharide S 30 moiety, a base B moiety, a purine or a pyrimidine and a signalling chemical moiety S_{ig} covalently attached thereto, either to the P, S or B moiety. Following are structural formulas of various base B moieties and 35 nucleotides which are modified in accordance with the practices of this invention.

The major purines

Adenine

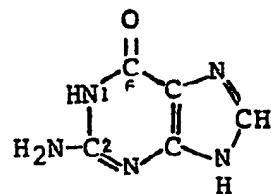
(6-aminopurine)

05



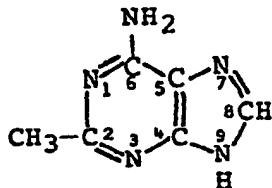
Guanine

(2-amino-6-oxypurine)

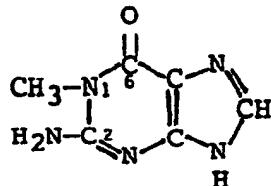
Two minor purines

10

2-Methyladenine



1-Methylguanine



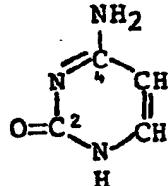
15

The major pyrimidines

Cytosine

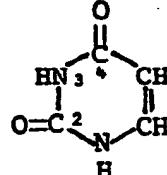
(2-oxy-4-aminopyrimidine)

20



Uracil

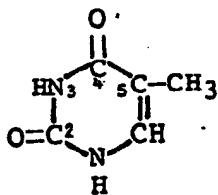
(2,4-dioxypyrimidine)



25

Thymine

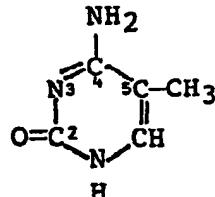
(5-methyl-2,4-dioxypyrimidine)



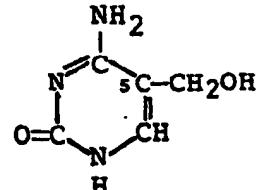
30

Two minor pyrimidines

5-Methylcytosine



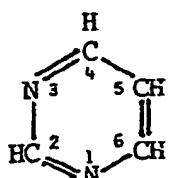
5-Hydroxymethylcytosine



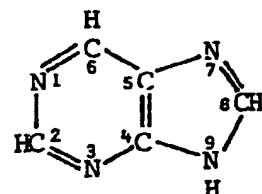
35

PYRIMIDINE

05



PURINE

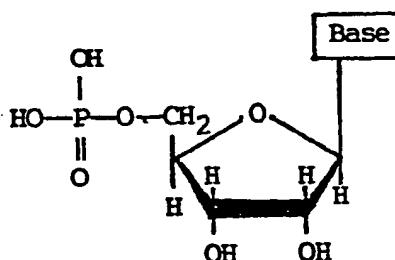
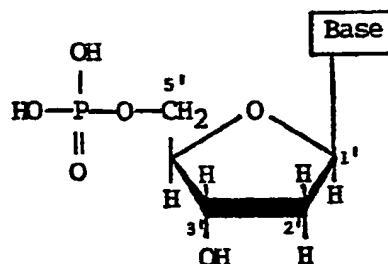


The major ribonucleotides and deoxyribonucleotides.

10

Ribonucleoside
5'-monophosphates

15

2'-Deoxyribonucleoside
5'-monophosphates

20

General Structure

General Structure

Names

Names

25

Adenosine 5'-phosphoric acid

(adenylic acid; AMP)

Guanosine 5'-phosphoric acid

(guanylic acid; GMP)

30

Cytidine 5'-phosphoric acid

(cytidylic acid; CMP)

Uridine 5'-phosphoric acid

(uridylic acid; UMP)

Deoxyadenosine 5'-phosphoric acid

(deoxyadenylic acid; dAMP)

Deoxyguanosine 5'-phosphoric acid

(deoxyguanylic acid; dGMP)

Deoxycytidine 5'-phosphoric acid

(deoxycytidylic acid; dCMP)

Deoxythymidine 5'-phosphoric acid

(deoxythymidylic acid; dTMP)

35

The special nucleotides in accordance with this invention, as indicated hereinabove, in addition to the P, S and B moieties, include a chemical moiety Sig covalently attached to the P, S and/or B moieties. Of 05 special interest in accordance with the practices of this invention would be those nucleotides having the general formula,

P - S - B - Sig
including mono-, di-, tri- or tetraphosphate,
wherein P is the phosphoric acid moiety, S the sugar
10 monosaccharide moiety, B the base moiety, either a purine or a pyrimidine. The phosphoric acid moiety P is attached at the 3' and/or the 5' position of the S moiety when the nucleotide is a deoxyribonucleotide and at the 2', 3' and/or 5' position when the nucleotide is 15 a ribonucleotide. The base B moiety is attached from the N1 position or the N9 position to the 1' position of the S moiety when the base moiety is a pyrimidine or a purine, respectively. The Sig moiety is covalently attached to the B moiety of the nucleotide and when so 20 attached is capable of signalling itself or makes it- self self-detecting or its presence known and desirably or preferably permits the incorporation of the resulting nucleotide P - S - B - Sig into or to form a double-stranded helical DNA or RNA or DNA-RNA hybrid 25 and/or to be detectable thereon.

Another special nucleotide in accordance with this invention is characterized by the general formula:

Sig

30

P - S - B

Such nucleotides in accordance with this invention would be characterized as ribonucleotides. The phosphoric acid moiety is attached at the 2', 3' and/or 5' position of the sugar S moiety and the base B being attached from the N1 position or the N9 position to the 1' position of the sugar S moiety when said base is a pyrimidine or a purine, respectively. The Sig chemical moiety is covalently attached to the sugar S moiety and said Sig chemical moiety when attached to said S moiety is capable of signalling itself or making itself self-detecting or its presence known and preferably permits the incorporation of the ribonucleotide into its corresponding double- stranded RNA or a DNA-RNA hybrid.

15 Sig

Such nucleotides P - S - B desirably have the Sig chemical moiety attached to the C2' position of the S moiety or the C3' position of the S moiety.

20 Still further, nucleotides in accordance with the practices of this invention include the nucleotides having the formula,

sig

25 **P = S = B**

wherein P is the phosphoric acid moiety, S the sugar moiety and B the base moiety. In these special nucleotides the P moiety is attached to the 3' and/or the 5' position of the S moiety when the nucleotide is deoxyribonucleotide and at the 2', 3' and/or 5' position when the nucleotide is a ribonucleotide. The base B is either a purine or a pyrimidine and the B moiety is attached from the N1 or the N9 position to the 1'

position of the sugar moiety when said B moiety is a pyrimidine or a purine, respectively. The Sig chemical moiety is covalently attached to the phosphoric acid P moiety via the chemical linkage

OH
·
- P - O - Sig
"
O ,
said Sig, when attached to said P moiety being capable
of signalling itself or making itself self-detecting or
its presence known and desirably the nucleotide is
capable of being incorporated into a double-stranded
polynucleotide, such as DNA, RNA or DNA-RNA hybrid and
when so incorporated therein is still self- detecting.

It is pointed out that the special nucleotides in accordance with the practices of this invention described or defined hereinabove by the general formula P - S - B - Sig, also include nucleotides wherein the Sig chemical moiety is covalently attached to the B moiety at the N⁶ or 6- amino group position when the B moiety is adenine or the N² or 2-amino group position when the B moiety is guanine or the N⁴ or 4-amino group position when the B moiety is cytosine. The resulting nucleotides containing the Sig moiety attached thereto are capable of signalling themselves or making themselves self-detecting or their presence known and being detectable is a double-stranded or DNA, RNA or DNA-RNA hybrid.

By way of summary, as indicated hereinabove with respect to the make-up of the various special nucleotides in accordance with this invention, the special nucleotides can be described as comprising a phosphoric acid moiety P, a sugar moiety S and a base moiety B, a purine or pyrimidine, which combination of P-S-B is well known with respect to and defines nucleotides, both deoxyribonucleotides and ribonucleotides. The nucleotides are then modified in accordance with the practices of this invention by having covalently attached thereto, to the P moiety and/or the S moiety and/or the B moiety, a chemical moiety Sig. The chemical moiety Sig so attached to the nucleotide P-S-B is capable of rendering or making the resulting nucleotide, now comprising P-S-B with the Sig moiety being attached to one or more of the other moieties, self-detecting or signalling itself or capable of making its presence known per se, when incorporated into a polynucleotide, especially a double-stranded polynucleotide, such as a double-stranded DNA, a double-stranded RNA or a double-stranded DNA-RNA hybrid. The Sig moiety desirably should not interfere with the capability of the nucleotide to form a double-stranded polynucleotide containing the special Sig-containing nucleotide in accordance with this invention and, when so incorporated therein, the Sig-containing nucleotide is capable of detection, localization or observation.

The Sig moiety employed in the make-up of the special nucleotides of this invention could comprise an enzyme or enzymic material, such as alkaline phosphatase, glucose oxidase, horseradish peroxidase or ribonuclease. The Sig moiety could also contain a fluorescing component, such as fluorescein or rhodamine or dansyl. If desired, the Sig moiety could include a

magnetic component associated or attached thereto, such as a magnetic oxide or magnetic iron oxide, which would make the nucleotide or polynucleotide containing such a magnetic-containing Sig moiety detectable by magnetic means. The Sig moiety might also include an electron dense component, such as ferritin, so as to be available by observation. The Sig moiety could also include a radioactive isotope component, such as radioactive cobalt, making the resulting nucleotide observable by radiation detecting means. The Sig moiety could also include a hapten component or per se be capable of complexing with an antibody specific thereto. Most usefully, the Sig moiety is a polysaccharide or oligosaccharide or monosaccharide, which is capable of complexing with or being attached to a sugar or polysaccharide binding protein, such as a lectin, e.g. Concanavilin A. The Sig component or moiety of the special nucleotides in accordance with this invention could also include a chemiluminescent component.

As indicated in accordance with the practices of this invention, the Sig component could comprise any chemical moiety which is attachable either directly or through a chemical linkage or linker arm to the nucleotide, such as to the base B component therein, or the sugar S component therein, or the phosphoric acid P component thereof.

The Sig component of the nucleotides in accordance with this invention and the nucleotides and polynucleotides incorporating the nucleotides of this invention containing the Sig component are equivalent to and useful for the same purposes as the nucleotides described in EP-A2-0 063 879.

More specifically, the

chemical moiety A described in EP-A2-0 063 879
is functionally the equivalent of
the Sig component or chemical moiety of the special
nucleotides of this invention. Accordingly, the Sig
component or chemical moiety of nucleotides of this
invention can be directly covalently attached to the P,
S or B moieties or attached thereto via a chemical
linkage or linkage arm as described in EP-A2-0 063 879

as indicated by the
dotted line connecting B and A of the nucleotides of
EP-A2-0 063 879. The various linker arms or
linkages identified in EP-A2-0 063 879 are
applicable to and useful in the preparation of the
special nucleotides of this invention.

A particularly important and useful aspect of the
special nucleotides of this invention is the use of
such nucleotides in the preparation of DNA or RNA
probes. Such probes would contain a nucleotide
sequence substantially matching the DNA or RNA sequence
of genetic material to be located and/or identified.
The probe would contain one or more of the special
nucleotides of this invention. A probe having a
desired nucleotide sequence, such as a single-stranded
polynucleotide, either DNA or RNA probe, would then be
brought into contact with DNA or RNA genetic material
to be identified. Upon the localization of the probe
and the formation of a double-stranded polynucleotide
containing the probe and the matching DNA or RNA
material to be identified, the resulting formed
double-stranded DNA or RNA-containing material would
then be observable and identified. A probe in
accordance with this invention may contain substantial-
ly any number of nucleotide units, from about 5 nucleo-
tides up to about 500 or more, as may be required. It
would appear that 12 matching, preferably consecutive,

nucleotide units would be sufficient to effect an identification of most of the DNA or RNA material to be investigated or identified, if the 12 nucleotide sequence of the probe matches a corresponding 05 cooperative sequence in the DNA or RNA material being investigated or to be identified. As indicated, such probes may contain one or more of the special Sig-containing nucleotides in accordance with this invention, preferably at least about one special 10 nucleotide per 5-10 of the nucleotides in the probe.

As indicated hereinabove, various techniques may be employed in the practices of this invention for the incorporation of the special nucleotides of this invention into DNA and related structures. One particularly useful technique referred to hereinabove involves the utilization of terminal transferase for the addition of biotinylated dUMP onto the 3' ends of a polypyrimidine or 15 to single-stranded DNA. The resulting product, such as a single-stranded or cloned DNA, which has biotinylated 20 dUMP attached to the 3' ends thereof, can be recovered by means of a Sepharose-avidin column wherein the avidin would complex with the biotinylated dUMP at the ends of 25 the DNA and be subsequently recovered. In accordance with the practices of this invention hybridization to mRNA could be accomplished in solution and the resulting hybrid recovered via a Sepharose-avidin column and the mRNA recovered therefrom. Similar techniques could be 30 employed to isolate DNA-RNA hybrids. This technique employing terminal transferase for the addition of the special nucleotides in accordance with this invention is widely applicable and the resulting modified nucleotides containing the special nucleotides in accordance with this invention including the special biotinylated nucleotides 35 or the special glycosylated nucleotides could be

1 selectively recovered via complexing with avidin upon a Sepharose-avidin column or complexing with a lectin, such as Concanavalin A or a Sepharose-Concanavalin A column.

5

Illustrative of the practices of this invention, biotinylated dUTP was added to the 3' ends of d pT 4 as well as single and double stranded DNA employing terminal transferase and the resulting product was purified through G-50 Sepharose and separated on a Sepharose-avidin affinity column. It was found that 69 % of the d pT 4 molecules were biotinylated and recovered on the Sepharose-avidin column. The results of this experiment established that terminal transferase added biotinylated dUMP to the 3' ends of a polypyrimidine.

10

15

20

25

30

35

The detection of nucleic acids to which specific molecules have been covalently attached can be effected through the use of many naturally occurring proteins to which small molecules are known to specifically bind.

05 In this procedure the small molecules are bound to the nucleotide using the allyl amine side chain. These nucleotides are then incorporated into specific nucleic acids using a DNA or RNA polymerase or ligase reaction or a chemical linkage. After annealing this probe with
10 a complementary antiprobe sequence, the presence of the probe is assayed for by the specific binding of the protein to the ligand covalently bound to the probe.

Examples of protein-ligand reactions that are
15 appropriate for this type of detector system include:

1. Enzymes and allosteric effector or modulator molecules. An example of this is the enzyme threonine dehydratase which is a heterotropic enzyme in that the
20 effector molecule, L-isoleucine, is different than the substrate, L-threonine, J. Monod, J. Wyman and J.P. Changeux (1965), J. Mol. Biol. 12:88-118.

2. Effector molecules involved in regulation. An example of this is the specific binding of 3',5-cyclic adenosine monophosphate to the cyclic AMP receptor
25 protein, I. Pastan and R. Perlman, Science 169:339-344 (1969). Another example is the lactose repressor molecule and the inducer molecules isopropylthiogalactoside or thiomethylgalactoside. These two inducer molecules are called gratuitous inducers in that they
30 are not metabolized by the enzymes they induce, W. Gilbert and B. Muller-Hill, Proc. Natl. Acad. Sci. (US), 70:3581-3584, (1973).

3. Hormone receptors and other receptors on the surface of the cell to which organic molecules will specifically bind. An example of this is the epinephrine-epinephrine receptor system in which
05 epinephrine is bound in a steriospecific manner with a high affinity to the receptor. With this system, since the receptor protein is insoluble in water, it will be imbedded in a lipid bilayer structure as for instance a liposome. Suitable detector systems would include
10 specific enzymes or fluorescent molecules inside or within the lipid bilayer.

4. Specific ligand binding proteins included in the transport of small molecules. An example of this is the periplasmic binding proteins of bacteria which have been
15 shown to bind many amino acids, glucose, galactose, ribose and other sugars, Pardee, A. Science, 162:632-637, (1968); G. L. Hazelbaur, and J. Adler, Nature New Bio. 230: 101-104, (1971).

20 In the above-mentioned examples the ligand bound to the nucleic acid reacts with a naturally occurring protein. The specificity of this reaction resides in the ligand-binding site of the protein.

25 One further example of small molecule interaction with naturally occurring proteins involves the specific binding of coenzyme or other prosthetic molecules to enzymes. Examples of such coenzymes include thiamin pyrophosphate, flavine mononucleotide, flavine adenine dinucleotide, nicotinamide adenine dinucleotide,
30 nicotinamide adenine dinucleotide phosphate, coenzyme A, pyridoxyl phosphate, biotin, tetrahydrofolic acid, coenzyme B₁₂, lipoic and ascorbic acid. Many of these molecules form covalent linkages with their apoenzymes.

However, some, for example, coenzyme A, coenzyme B₁₂ and tetrahydrofolic acid, associate in a non-covalent but specific manner with their cognate apoenzymes. A specific coenzyme-apoenzyme system for use in this system is flavine adenine dinucleotide (FAD) and flavine 05 adenine dinucleotide reductase isolated from Escherichia coli. With this system the binding of FAD is sufficiently strong to permit detection.

The special nucleotides of this invention and poly- 10 nucleotides including such nucleotides, either single-stranded or double-stranded polynucleotides, DNA and/or RNA, comprising the components, phosphoric acid moiety P, the sugar or monosaccharide moiety S, the base moiety B, a purine or a pyrimidine, and the signalling or 15 self-detecting moiety, Sig, covalently attached to either the P, S or B moieties, as indicated hereinabove, have many uses and utilities. For example, the nucleotides of this invention and polynucleotides containing the nucleotides of this invention are useful 20 as immune-stimulating agents, as adjuvants in vaccines, as agents for the stimulation or induction from competent cells, such as lymphocytes, for the production of lymphokines, cytokines or cytokinins, interferon or other cellular products.

25 It is well known that double-stranded poly A:U is a stimulator or inducer for the production of interferon, although weakly so. Similarly, poly I:C is also known as a stimulator or inducing agent for the production of 30 interferon.

The advantage of polynucleotides, such as double-stranded polynucleotides incorporating one or more nucleotides in accordance with this invention is that, in effect, such

polynucleotides would be more effective and more powerful inducing or stimulating agents for the production of interferon and related materials from cells. For example, nucleotides in accordance with this
05 invention containing the above-described components P, S, B and Sig, are suitably prepared so that the nucleotides and polynucleotides prepared therefrom are more resistant to nucleases. Similarly, such nucleotides and polynucleotides containing the same and
10 suitably prepared which are more capable of contacting, stimulating and penetrating cellular surfaces or membranes, such as the cellular surfaces or membranes of lymphocytes and other cells so as to stimulate the same for the production of a desired cellular product, such
15 as interferon.

Particularly useful among those special nucleotides in accordance with this invention having the formula P-S-B-Sig and especially useful are those wherein the Sig
20 component is at the 5 position of the pyrimidine or the 7 position of the purine or a deazapurine or the N² position of guanine or the N⁶ position of adenine. Such nucleotides and polynucleotides incorporating the same, both single-stranded and double-stranded nucleotides,
25 DNA and/or RNA are prepared in accordance with this invention to provide increased stability with respect to the double-stranded helix of DNA or RNA or DNA-RNA hybrid containing the same. Increased resistance to nucleases is also achievable as well as alterations or
30 favorable changes in the hydrophobic properties or electrical or charge properties of the nucleotides and polynucleotides containing the same. Also, nucleotides and polynucleotides in accordance with this invention are prepared which, when administered to humans, have

reduced pyrogenicity or exhibit less other whole body toxic responses. Additionally, the nucleotides and polynucleotides are prepared in accordance with this invention to provide a ligand, such as the component

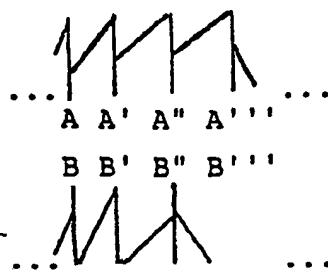
05 Sig, to which specific polypeptides can combine to create or bring about the formation of RNA complexes. It is seen therefore that the nucleotides of this invention include the P, S, B and Sig components wherein the Sig is covalently attached to either the P, S or B
10 moieties open up or provide a whole array of chemical agents having special biological effects including therapeutic effects and cytotoxic effects.

The special nucleotides of this invention, including
15 polynucleotides containing these nucleotides, in addition to being stimulating or inducing agents for the production of cellular materials or products, such as interferons, lymphokines and/or cytokines, are also useful for their chemotherapeutic effect and for the
20 preparation of chemotherapeutic agents based thereon but also for their cytotoxic effects and the production of cytotoxic agents based thereon. The moiety Sig attached to the special nucleotides of this invention containing the other moieties or components P, S, B
25 provides a site per se for the attachment thereto, the Sig component, of special agents of known chemotherapeutic or cytotoxic effect. Such nucleotides could be introduced or administered to the subject being treated, e.g. human body or animal, so as to be
30 incorporated into the DNA and/or RNA components of the body or cell so as to either interfere with the synthesis of the body or cellular DNA and/or RNA or to attack tumors or to, in effect, kill or otherwise interfere with the growth of undesired cells.

The administration of the nucleotides and/or poly-nucleotides containing the nucleotides to the body, human body or animal, can be effected by a number of suitable means. Particularly effective would be the
05 intravenous introduction to the body of preparations containing the nucleotides of this invention and a suitable physiologically acceptable carrier or the nucleotides could be administered subcutaneously, transdermally, or intramuscularly or by direct intro-
10 duction into the site where the chemotherapeutic or cytotoxic effect of the nucleotides is sought or desired to be effective. Not only could desired chemotherapeutic or cytotoxic effects be achieved systemically or locally but also, as indicated
15 hereinabove, the special P, S, B and Sig-containing nucleotides of this invention, including polynucleo- tides containing such nucleotides, are useful as immune-stimulating agents and adjuvants therefor. Accordingly, vaccines containing the special nucleo-
20 tides and polynucleotides in accordance with this invention can be prepared having improved effectiveness and versatility.

Of special interest in the practices of this invention
25 improved polynucleotides incorporating the special nucleotides of this invention are provided as inducers or stimulating agents for the production of interferon. Such polynucleotides would be single-stranded or double-stranded ribonucleotides, dsRNA, having the
30 structures,

05



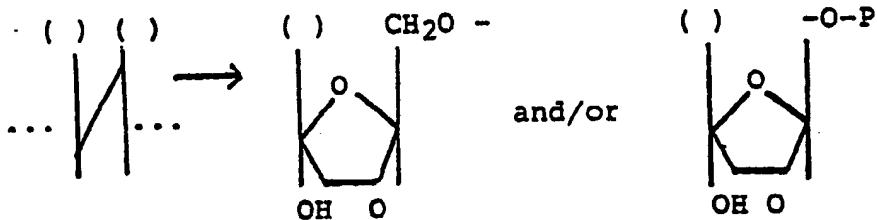
10

or



where A and B are complementary base pairs, such as a purine, a 7-deazapurine or pyrimidine modified by the addition of an organic moiety Sig in accordance with
15 the disclosures of this invention on the 5 position of the pyrimidine ring or the 7 position of the purine ring or the N² of guanine, or the N⁶ of adenine or the N⁴ of cytosine as described herein. The modifications of the polynucleo-
20 tides at these positions lead to relatively undisrup- tive or non-disruptive double-stranded nucleic acid molecules as measured by rates of association and melting points. In the special polynucleotides of this invention employed as inducers of interferon and other
25 cellular or humoral factors or components, such as lymphokines or cytokines, the following groups would be attached thereto as indicated by the formulas,

30



35

In the utilization of the special polynucleotides of this invention, such as the special dsRNA of this invention in the induction process for the production of interferon it has been demonstrated that DEAE-dextran facilitates this operation. It appears that since DEAE-dextran complexes with dsRNA and protects it for nuclease degradation, thereby enhancing interferon induction. It has also been noted that poly rC : rI is taken into cells more efficiently when complexed with DEAE-dextran. Accordingly, in the practices of this invention the hydrophobic properties and the ionic or electron charge properties of the special dsRNA of this invention are important factors and capable of manipulation in the applicability of these materials to induce interferon production. It has been observed that such conditions or factors which promote the induction of interferon also lead to and promote the induction of other cellular or humoral components, such as lymphokines and cytokines. It is apparent, therefore, that the special nucleotides and polynucleotides containing the special nucleotides of this invention act as immune modulators and stimulators of the immune response other than simply being effective as inducers of interferon production. Superior agents for the above in accordance with the practices of this invention would include nucleotides wherein the Sig moiety incorporates biotin or streptavidin or avidin.

Poly rI:poly rC complexed poly L-lysine exhibits adjuvant properties and such properties are enhanced and improved in accordance with the practices of this invention when the poly rI and poly rC components are modified to include one or more of the special nucleotides in accordance with this invention.

The preparation of DNA probes in accordance with another aspect of this invention can be carried out in a manner which does not require the preparation or utilization of the special nucleotides described
05 herein. For example, double-stranded DNA can be reacted with a carcinogen or alkylating agent. After the carcinogen has reacted with or alkylated the double-stranded DNA, the resulting modified DNA is melted to produce a DNA hybridizing probe containing the reaction product of the DNA and the carcinogen or alkylating agent. When thus-modified or reacted DNA is employed as a hybridizing probe, any resulting formed double helix or double-stranded DNA would be assayed or searched out by means of a double antibody technique.
10 The primary antibody would be an anti-carcinogen and the secondary antibody would be horseradish-peroxidase conjugated anti-peroxidase antibody. The advantage of this technique is that it would be easy to label the double-stranded DNA. This special approach is indicated hereinabove in the examples accompanying the
15 description of this invention and is generally applicable for the preparation of DNA probes from double-stranded or double helical DNA. However, this procedure is a disruptive procedure involving the modification of the double helical deoxyribonucleotide
20 polymer or DNA.
25

In the description of the special nucleotides and modified DNA employed or developed in the practices of this invention, mention has been made of mono, oligo and polysaccharides. It is pointed out that derivatives of mono, oligo and polysaccharides are also useful in the preparation of the special nucleotides of this invention. For example, it is possible to modify
30 individual sugar moieties employed in the make-up of
35

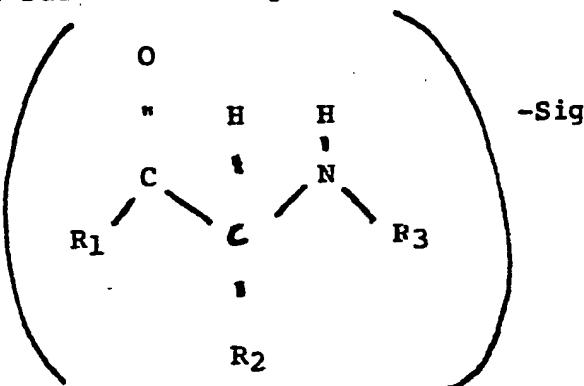
the special nucleotides and employ the resulting modified sugar moieties to effect or carry out additional chemical reactions. Such modified mono, oligo and polysaccharide moieties, when employed as the
05 Sig moiety in the preparation of the special nucleotides of this invention, provide an added versatility with respect to the detection of the nucleotides or other compounds containing such modified saccharides either as the sugar S or as the Sig moiety thereof.

10

In another aspect of this invention the Sig moiety instead of being attached to a nucleotide could also be attached to proteins. Not only could such proteins be attached to nucleotides or polynucleotides but also
15 such proteins could be identified per se whether attached to a nucleotide or polynucleotide or unattached. In accordance with the practices of this aspect of the invention, a suitable such protein adduct would have the formula,

20

25



30

wherein R₁ is an OH or an amino acid or acids and R₂ is an amino acid side chain and R₃ is H or an amino acid or acids and Sig is attached to the R₁ and/or R₂ and/or R₃.

1 WHAT IS CLAIMED IS:

1. A nucleotide having the general formula P-S-B-Sig wherein P is the phosphoric acid moiety, S the sugar or monosaccharide moiety, B being the base moiety, the phosphoric acid moiety being attached at the 3' and/or the 5' position of the sugar moiety when said nucleotide is a deoxyribonucleotide and at the 2', 3' and/or 5' position when said nucleotide is a ribonucleotide, said base being a purine or a pyrimidine, said base being attached from the N1 position or the N9 position to the 1' position of the sugar moiety when said base is a pyrimidine or a purine, respectively, and wherein said Sig is a chemical moiety covalently attached to the base B of said nucleotide, said Sig when attached to said base B being capable of signalling itself or makes itself self-detecting or its presence known.
2. A nucleotide in accordance with Claim 1 wherein Sig is attached to B at a position such that said nucleotide is capable of being incorporated into or to form a double-stranded ribonucleic acid, a double-stranded deoxyribonucleid acid or a double-stranded deoxyribonucleic acid-ribonucleic acid hybrid.
3. A nucleotide in accordance with Claim 1 wherein said Sig chemical moiety is a sugar selected from the group consisting of triose or tetrose, or pentose, a hexose, or heptose, and an octose.
4. A nucleotide in accordance with Claim 1 wherein said Sig chemical moiety comprises a component selected from the group consisting of an electron dense component, a magnetic component, an enzyme, a hormone component, a radioactive component, a metal-containing component, a fluorescing component and an antigen or antibody component.

1 5. A nucleotide in accordance with Claim 1 wherein
said Sig chemical moiety is a sugar residue and wherein
said sugar is complexed with or attached to a sugar or
polysaccharide binding protein.

5

6. A nucleotide in accordance with Claim 1 wherein said
Sig chemical moiety is connected to said base B via a
chemical linkage.

10

7. A nucleotide in accordance with Claim 6 wherein said
chemical linkage includes an olefinic bond at the a-
position relative to base B.

15

8. A nucleotide in accordance with Claim 6 wherein said
chemical linkage includes the moiety,

- CH₂ - NH - .

20

9. A nucleotide in accordance with Claim 1 wherein said
Sig chemical moiety includes or comprises a catalytic
metal component.

25

10. A single-stranded or a double-stranded polynucleotide
comprising one or more nucleotides in accordance with
Claim 1.

11. A ribonucleotide having the general formula,

Sig

P - S - B

30

wherein P is the phosphoric acid moiety, S the sugar
moiety and B the base moiety, the phosphoric acid moiety
being attached at the 2', 3' and/or 5' position of the
sugar moiety, said base B being attached from the N1
position or the N9 position to the 1' position of the
sugar moiety when said base is a pyrimidine or a purine,

1 respectively, and wherein said Sig is a chemical moiety
covalently attached to the sugar S, said Sig, when
attached to said sugar S, being capable of signalling
itself or making itself self-detecting or its presence
5 known.

12. A ribonucleotide in accordance with Claim 11 wherein
said Sig chemical moiety includes or comprises an enzyme.

10 13. A polyribonucleotide coupled or attached to a poly-
peptide.

14. A nucleotide having the general formula

Sig

15

' P - S - B

wherein P is the phosphoric acid moiety, S the sugar
moiety and B the base moiety, the phosphoric acid moiety
being attached to the 3' and/or the 5' position of the
20 sugar moiety when said nucleotide is a deoxyribonucleotide
and at the 2', 3' and/or 5' position when said nucleotide
is a ribonucleotide, said base B being a purine or
pyrimidine, said base B being attached from the N1
position or the N9 position to the 1' position of the
25 sugar moiety when said base B is a pyrimidine or a purine,
respectively, and wherein Sig is a chemical moiety
covalently attached to the phosphoric acid moiety via the
chemical linkage

OH

30

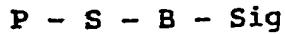
' - P - O - Sig,

"

O

said Sig, when attached to said phosphoric acid moiety P
85 being capable of signalling itself or making itself self-
detecting or its presence known.

1 15. A nucleotide having the general formula



wherein P is the phosphoric acid moiety, S the sugar and monosaccharide moiety, B being the base moiety, the phosphoric acid moiety being attached to the 3' and/or the 5' position of the sugar moiety when said nucleotide is deoxyribonucleotide and at the 2', 3' and/or 5' position when said nucleotide is a ribonucleotide, said base being a purine or a pyrimidine, said base being attached from the N1 position or the N9 position to the C1' position of the sugar moiety when said base is a pyrimidine or a purine, respectively, and wherein said Sig is a chemical moiety covalently attached to the base B of said nucleotide, said Sig being attached to the N⁶ or 6-amino group when said base B is adenine or the N² or 2-amino group when said base B is guanine or the N⁴ or 4-amino group when said base B is cytosine, said Sig when attached to said base B being capable of signally itself or makes itself self-detecting or its presence known.

16. A nucleotide having the general formula P-S-B, wherein P is the phosphoric acid moiety, S the sugar or monosaccharide moiety and B the base moiety, said nucleotide having covalently attached to the P or S or B moiety a chemical moiety Sig, said Sig chemical moiety when attached to the P moiety is attached thereto via the chemical linkage,

30



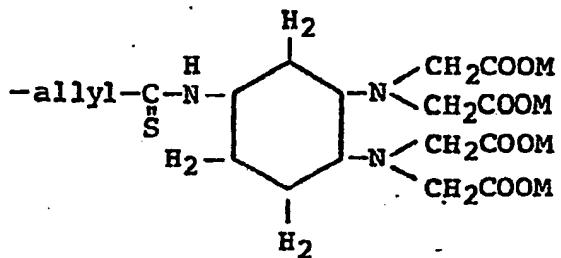
35 and when Sig is attached to the S moiety, the S moiety is a ribose group, said chemical moiety Sig when attached to said P, S or B being capable of signalling itself or makes itself self-detecting or its presence known.

1 17.. A polynucleotide comprising one or more nucleotides
in accordance with Claim 1 or Claim 11 or Claim 14 or
Claim 15 or Claim 16, coupled or attached to a poly-
peptide, said polypeptide having attached thereto one or
5 more Sig chemical moieties, said Sig when attached to said
polypeptide being capable of signalling itself or making
itself self-detecting or its presence known.

18. A nucleotide in accordance with Claim 1 or Claim 11
10 or Claim 14 or Claim 15 or Claim 16 wherein said Sig chemical
moiety includes a chelating agent.

19. A nucleotide in accordance with Claim 1 or Claim 11
or Claim 14 or Claim 15 or Claim 16 wherein the Sig chemical
15 moiety comprises the chemical moiety

20



20. A nucleotide in accordance with Claim 1, 11, 14, 15 or
16 wherein said Sig chemical moiety includes a coenzyme.

25

21. A polynucleotide containing one or more nucleotides
in accordance with Claim 1, or Claim 11, or Claim 14, or
Claim 15, or Claim 16 wherein said Sig moiety is radio-
30actively labeled.

22. A polynucleotide containing one or more nucleotides
in accordance with Claims 11, 14, 15 or 16 wherein said Sig
chemical moiety comprises biotin attached to radioactively
35labeled avidin, or to radioactively labeled strepavidin.

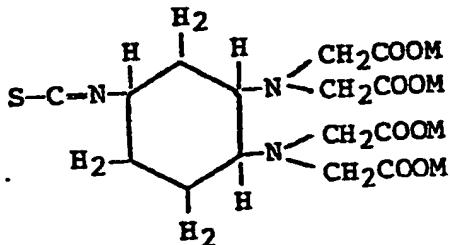
1 23. A polynucleotide containing one or more nucleotides,
 in accordance with Claim 1 or Claim 11, or Claim 14, or
 Claim 15, or Claim 16, wherein said Sig moiety comprises
 a radioactively labeled antibody or a radioactively
 5 labeled protein.

24. A method for the stimulation or induction of cells
 for the production of lymphokines, cytokinins and/or
 interferon which comprises introducing into or bringing
 10 into contact with cells capable of and/or functioning for
 the production of said lymphokines, cytokinins and/or
 interferon an effective lymphokine, cytokinin and/or
 interferon stimulating and production inducing amount of a
 nucleotide in accordance with Claim 1, 11, 14, 15, or 16.

15

25. The compound

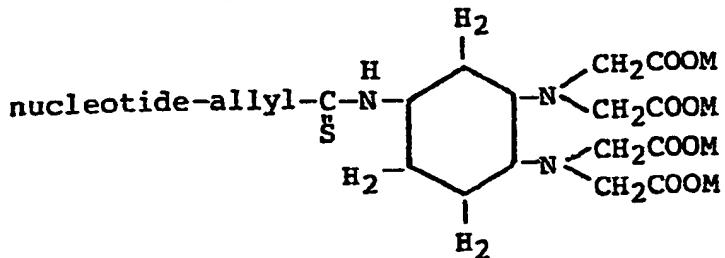
20



25 wherein M is hydrogen or a metal.

26. The compound

30

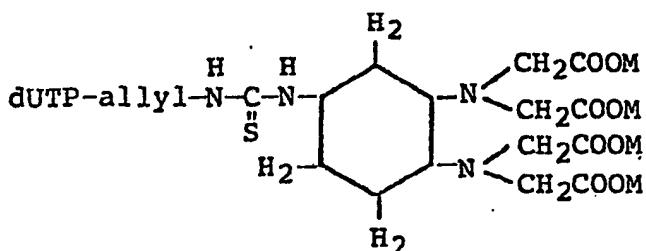


35

wherein M is hydrogen or a metal.

1 : 27. The compound

5



wherein M is hydrogen or a metal.

10.

28. An amino acid or polypeptide comprising a Sig moiety attached thereto.

15

29. A monosaccharide or a polysaccharide comprising a Sig moiety attached thereto.

20

30. A method of detecting a first compound which includes a nucleotide in accordance with Claim 1, 11, 14, 15 or 16 as part of said first compound, which comprises contacting said first compound with a second compound capable of forming a complex therewith under suitable conditions so as to form said complex, said complex comprising said first compound and said second compound and detecting said complex.

25

31. A method of determining the presence of a deoxyribonucleic or ribonucleic acid molecule which comprises forming a double-stranded hybrid polynucleotide duplex which includes a single strand of deoxyribonucleic acid or ribonucleic acid corresponding to or derived from said deoxyribonucleic or ribonucleic acid molecule and a nucleotide in accordance with Claim 1, 11, 14, 15 or 16 and detecting said duplex.

32. A method of testing a bacterium to determine its resistance to an antibiotic which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said bacterium which confers resistance of said bacterium to said antibiotic and which includes a nucleotide in accordance with Claim 1,11,14,15 or 16 incorporated therein, contacting said polynucleotide under suitable conditions with a deoxyribonucleic acid obtained from said bacterium so as to form a double-stranded hybrid duplex and detecting the presence of said duplex, the detection of said duplex indicating resistance of said bacterium to said antibiotic and the absence of said duplex indicating the susceptibility of said bacterium to an antibiotic. --

33. A method of diagnosing a genetic disorder in a subject which comprises preparing a polynucleotide complementary to the deoxyribonucleic acid gene sequence of said subject which is associated with said genetic disorder and includes a compound in accordance with Claim 1,11,14,15 or 16, incorporated therein contacting said polynucleotide under suitable conditions with deoxyribonucleic acid obtained from said subject so as to form a double-stranded hybrid duplex and detecting the presence of said hybrid duplex, the presence or absence of said hybrid duplex indicating the presence or absence of said genetic disorder. --

1 34. A method of diagnosing thalassemia which comprises preparing
a polynucleotide complementary to the deoxyribonucleic
acid gene sequence which is absent in β -minus-thalassemia
subjects and includes a nucleotide in accordance with
5 Claim 1, 11, 14, 15 or 16, contacting said polynucleotide
under suitable conditions with deoxyribonucleic acid
obtained from said subject so as to form a double-stranded
hybrid duplex and determining the presence of said duplex,
the absence of said duplex indicating the presence of β -
10 minus-thalassemia.

15 35. A method of chromosomal karyotyping which comprises
preparing a series of modified polynucleotides correspond-
ing to a series of defined genetic sequences located on
chromosomes, said polynucleotides including one or more
compounds in accordance with Claim 1, 11, 14, 15 or 16,
contacting said polynucleotides with deoxyribonucleic
acid of or obtained from chromosomes so as to form hybrid
duplexes and detecting said duplexes, thereby determining
20 the location of said duplexes on said chromosomes and the
location of said genetic sequences on said chromosomes.

25 36. A method of diagnosing a tumor cell which comprises
preparing a polynucleotide which is complementary to a
messenger ribonucleic acid synthesized from a deoxy-
ribonucleic acid gene sequence associated with the pro-
duction of a polypeptide diagnostic for or identifiable
with said tumor cell or which is complementary to said
deoxyribonucleic acid gene sequence and which includes
30 a compound in accordance with Claim 1, 11, 14, 15 or 16
introducing said polynucleotide into contact with said
cell under suitable conditions so as to permit said poly-
nucleotide to hybridize with said deoxyribonucleic acid
gen sequence or said messenger ribonucleic acid and
35 detecting the formation of a resulting formed hybrid
containing said polynucleotide.

37. A method of determining or identifying or diagnosing a first compound capable of complexing with or binding with a first component of a second compound, said second compound comprising said first component and a second component, said second component being attached to or complexed with said first component, which comprises bringing said first compound into contact with said second compound whereby said first component of said second compound complexes with said first compound to form a third compound comprising said first compound and said second compound and contacting said resulting third compound with a fourth compound capable of binding with or attaching to said second component of said second compound making up said third compound. --

38. A method of determining or identifying or diagnosing a first compound capable of complexing with or binding with an amino acid, peptide or protein which comprises bringing said first compound into contact with a second compound, said second compound comprising an amino acid, peptide or protein attached to or complexed with a sugar, or oligosaccharide or polysaccharide, whereupon said second compound complexes with or becomes attached to said first compound to form a third compound comprising said first compound and said second compound and contacting said resulting third compound with a protein or amino acid or peptide capable of binding with or attaching to the sugar or oligosaccharide moiety of said second compound making up said third compound. --

39. A method of determining or identifying or diagnosing a first compound capable of complexing or binding with a first component of a second compound, said second compound comprising said first component, a second component attached to or complexed with said first component, said second compound being a chelating agent, and a third component, said third component being an ion chelated or fixed to said second component, which comprises bringing said first compound into contact with said second compound whereby said first component of said compound complexes with said first compound to form a third compound comprising said first compound and said second compound.

40. A method of determining or identifying or diagnosing in an organism, cellular material or tissue a first compound comprising a receptor molecule with a receptor site that specifically binds to a second molecule which comprises bringing said organism, material or tissue containing said receptor molecule into contact with a second compound, said second compound comprising a first component capable of complexing with or binding with said receptor molecule, said second compound also comprising a second component, said second component being attached to or complexed with said first component, said first component of said second compound when in contact with said receptor molecule complexes with said receptor molecule to form a third compound comprising said first compound and said second

compound and contacting said third compound with a fourth compound capable of binding with or attaching to said second component of said second compound making up said third compound, said first component of said second compound being selected from group A consisting of an amino acid, a peptide, or a protein or from group B consisting of a sugar, oligosaccharide or polysaccharide or from group C consisting of a purine, pyrimidine, nucleoside, nucleotide, oligonucleotide or polynucleotide, said second component of said second compound being a sugar, oligosaccharide or polysaccharide and said fourth compound is selected from the group consisting of a peptide or polypeptide, a lectin, or an antibody. --

41. A method of detecting a first compound which includes in its make-up a nucleotide in accordance with Claim 20, which comprises contacting said first compound with an apoenzyme corresponding to said coenzyme. --

42. A method of determining the presence of a polynucleotide, comprising one or more nucleotides in accordance with Claim 1, 11, 14, 15 or 16, wherein said Sig component of said nucleotide comprises biotin which comprises bringing said polynucleotide into contact with radioactive labeled avidin or strepavidin to bind said biotin to said radioactive labeled avidin or strepavidin and determining the presence of the biotin bound to said radioactive labeled avidin or strepavidin by detecting the radioactivity of the biotin-bound radioactive avidin or strepavidin.

1 43. A diagnostic kit useful for determining the presence
of a nucleic acid-containing organism or the like which
comprises a polynucleotide which includes in its make-up
a compound selected from the compounds of Claim 1, 11, 14,
5 15 or 16 and which is complementary to all or an identifiable,
distinct or unique portion of the nucleic acid contained
in said organism and means for detecting or expressing
the presence or absence of a resulting formed hybrid bet-
ween said polynucleotide and said nucleic acid of said
10 organism when said polynucleotide is brought into contact
with the nucleic acid of said organism or the like under
hybrid forming conditions.

15

20

25

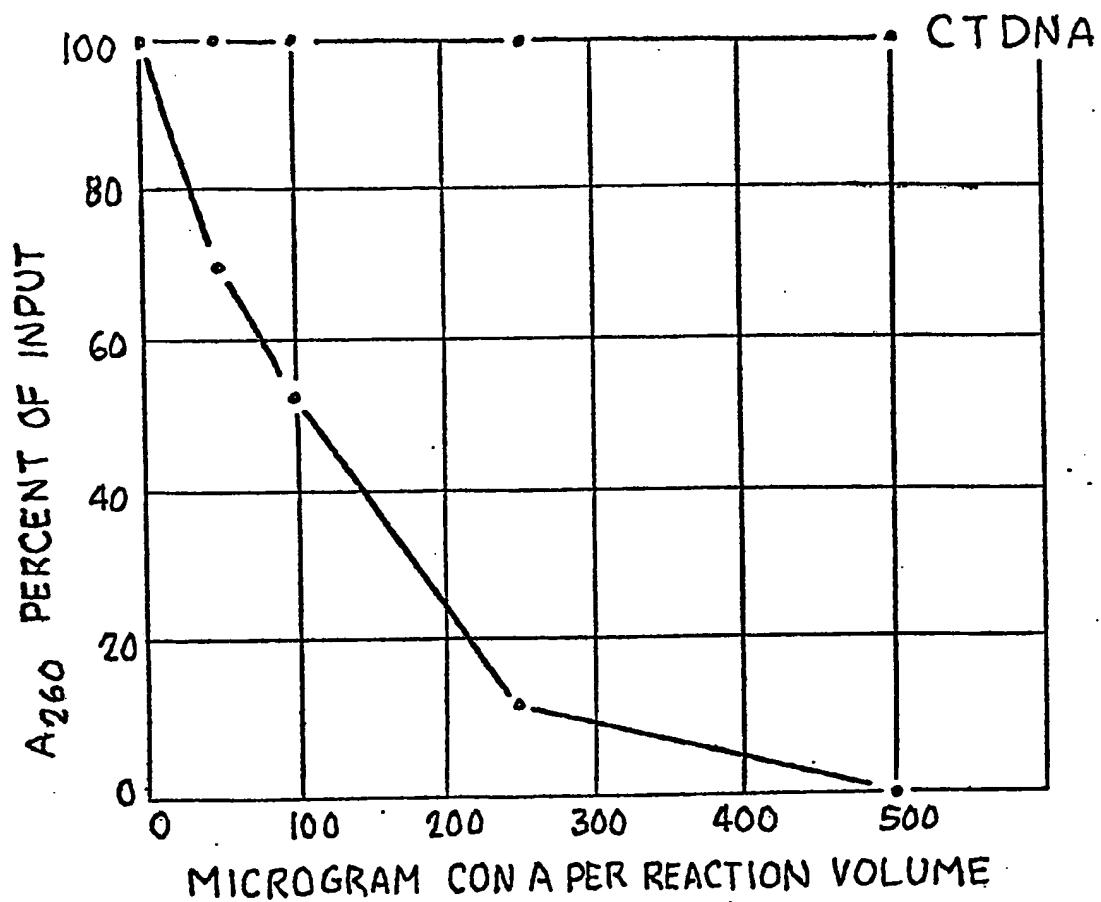
30

35

0097373

1/4

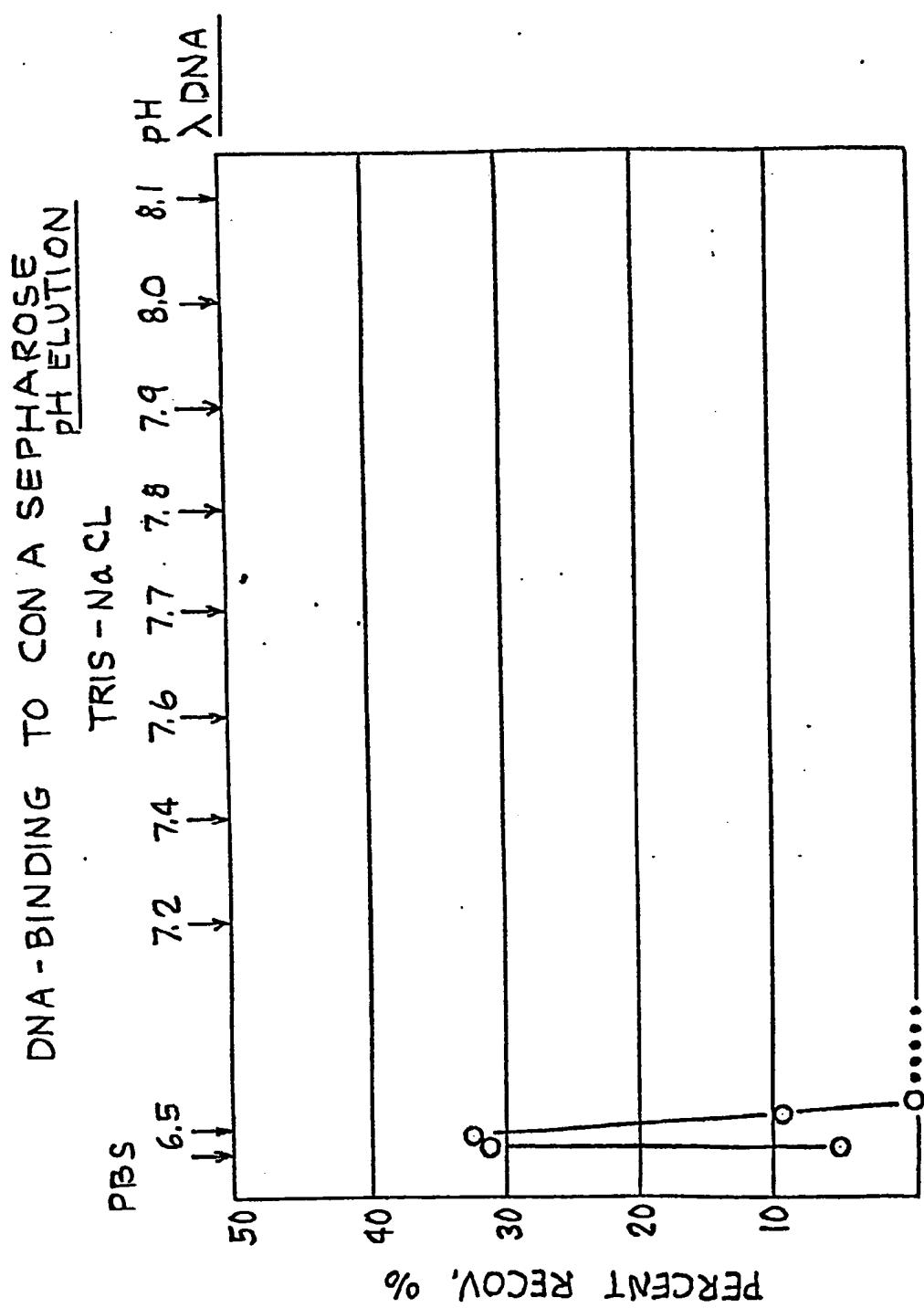
Fig. 1.



0097373

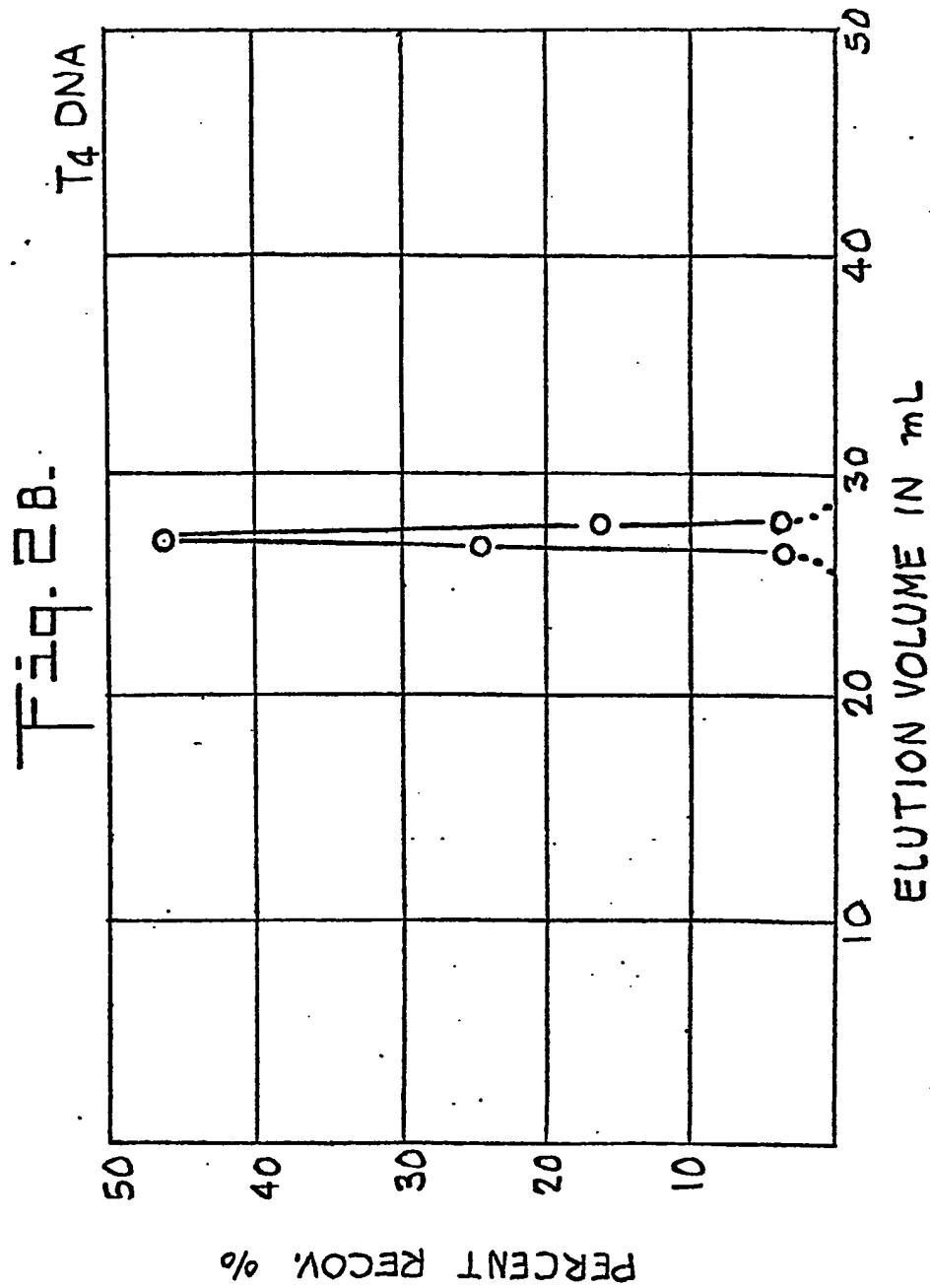
2/4

Fig. 2A.



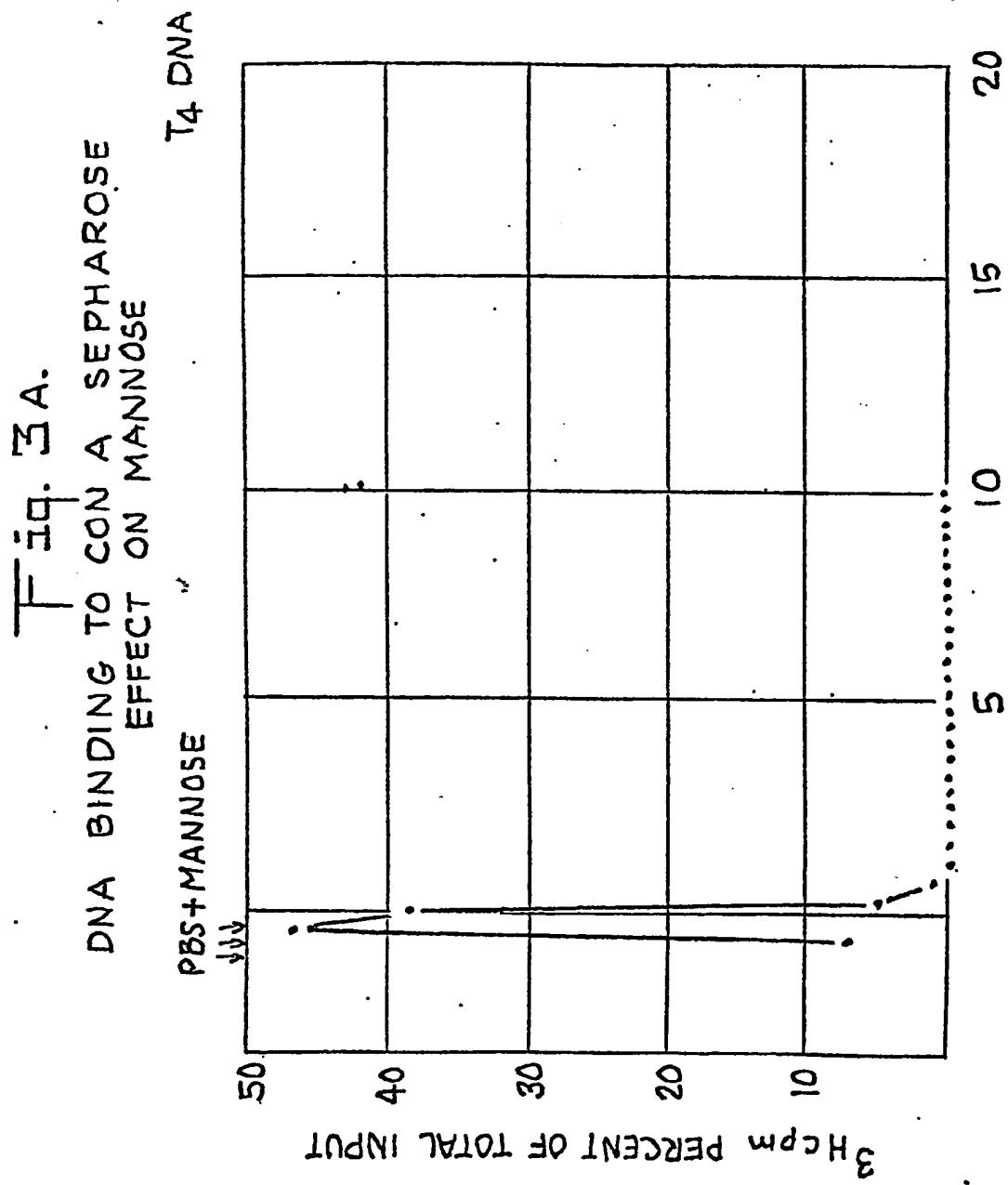
0097373

73/4



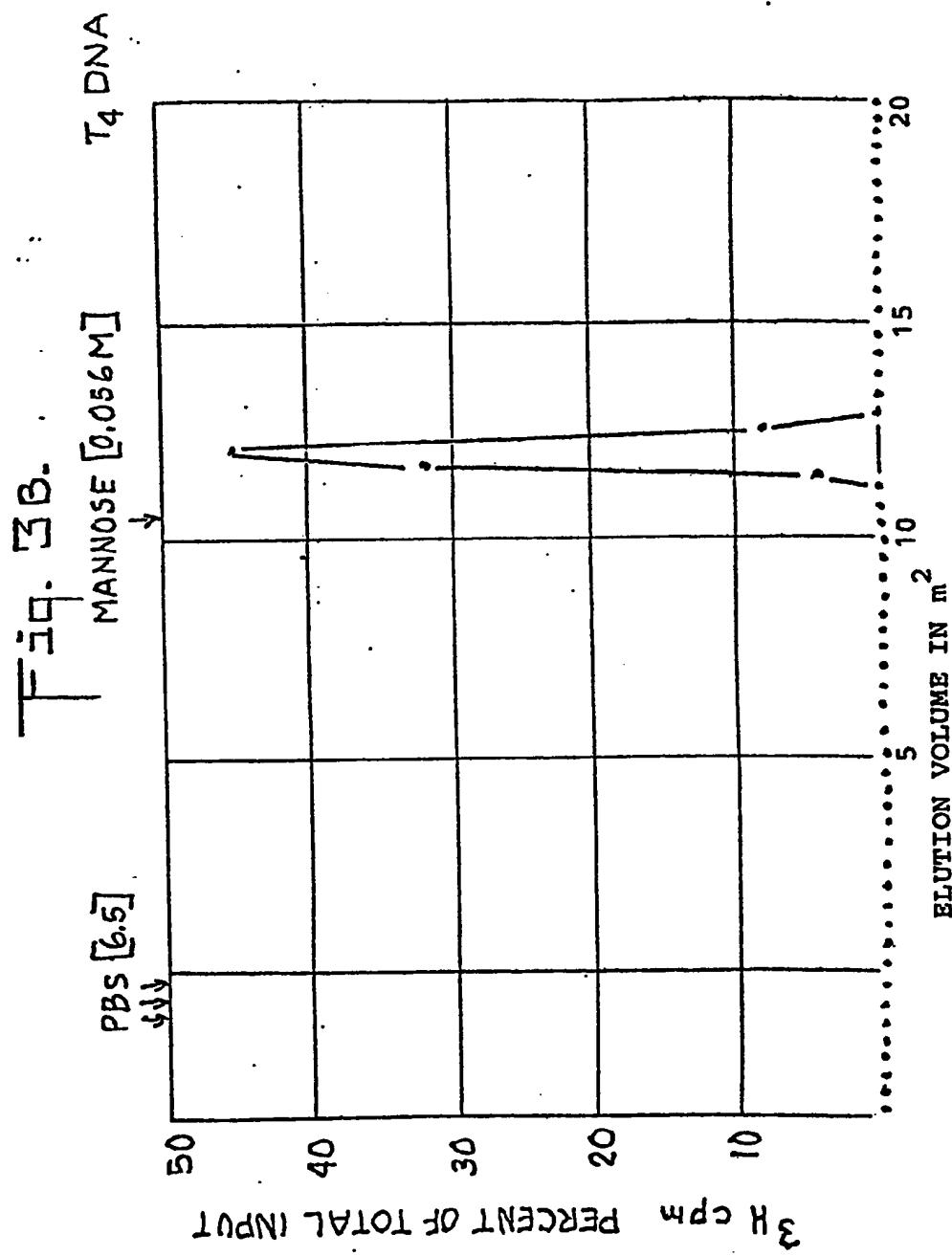
0097373

5/
x



0097373

5/
Y



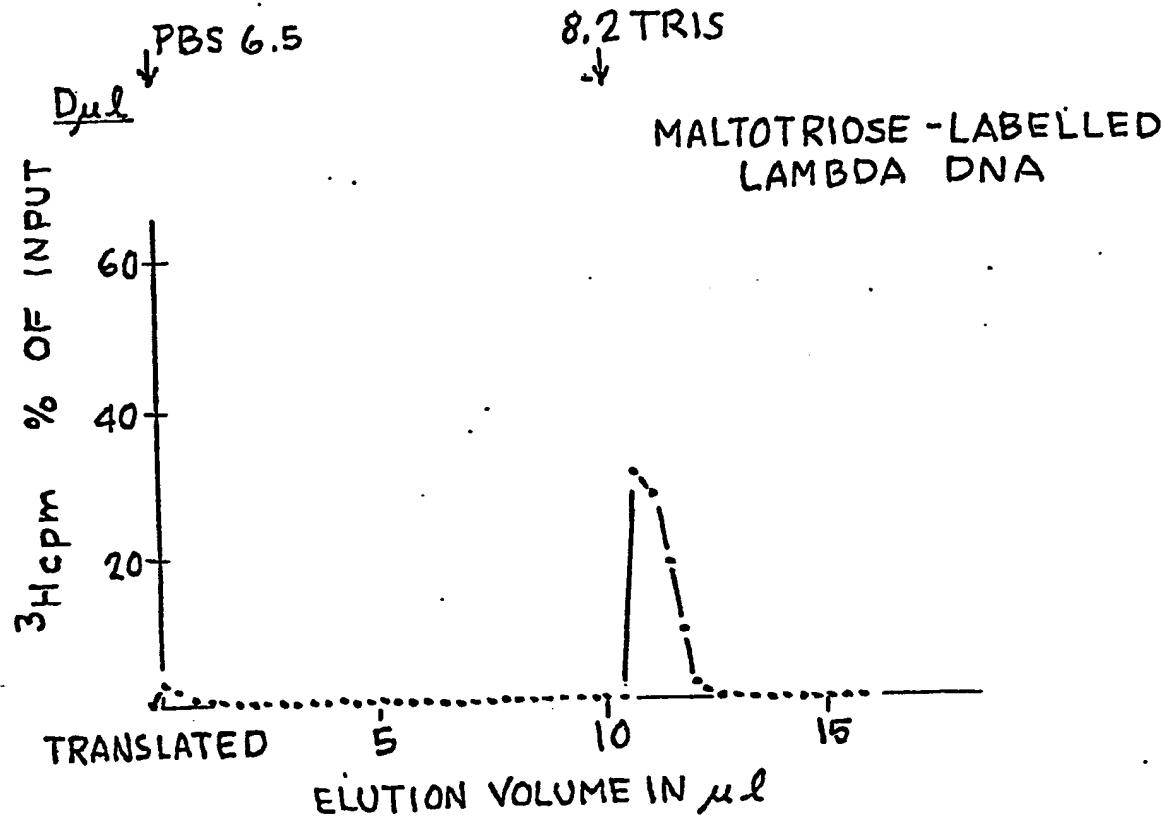
- 12 -

0097373

6/4

Fig. 4A.

CON-A SEPHAROSE BINDING OF GLUCOSYLATED DNAs
3/8 - 9/82



0097373

4/4

Fig. 4B.

